Copy 263 RM 51B05

NACA RM 51B05

6335

Chassificative

NACA



RESEARCH MEMORANDUM

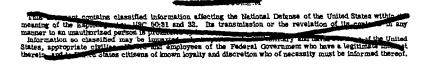
PROPERTIES OF LOW-CARBON N-155 ALLOY BAR STOCK

FROM 1200° TO 1800° F

By J. W. Freeman and A. E. White

University of Michigan

AFMDC TECKNOWL LODGEN AFL 2811



NATIONAL ADVISORY COMMITTEE FOR AERONAUTICS

WASHINGTON May 3, 1951

RESTRICTED

319.98/13

National Aeronautics and Space Administration

Langley Research Center Hampton, Virginia



Recty to Alic of

139A

JUN 1 6 1983

TO:

Distribution

FROM:

180A/Security Classification Officer

SUBJECT:

Authority to Declassify NACA/NASA Documents Dated Prior to

January 1, 1960

(informal, carrespondence)

Effective this date, all material <u>classified by this Center</u> prior to January 1, 1960, is declassified. This action does not include material derivatively classified at the Center upon instructions from other Agencies.

Immediate re-marking is not required; however, until material is re-marked by lining through the classification and annotating with the following statement, it must continue to be protected as if classified:

"Declassified by authority of LaRC Security Classification Officer (SCO) letter dated June 16, 1983," and the signature of person performing the re-marking.

If re-marking a large amount of material is desirable, but unduly burdensome, custodians may follow the instructions contained in NRB 1640.4, subpart F, section 1203.604, paragraph (h).

This declassification action complements earlier actions by the National Archives and Records Service (NARS) and by the NASA Security Classification Officer (SCO). In Declassification Review Program 807008, NARS declassified the Center's "Research Authorization" files, which contain reports, Research Authorizations, correspondence, photographs, and other documentation. Barlier, in a 1971 letter, the NASA SCO declassified all NACA/NASA formal series documents with the exception of the following reports, which must remain classified:

First Author

E-51A30 E-53G20 E-53G21 E-53K18 BL-54J21a

Nagey Francisco Johnson Spooner Westphal Fox

FOX Himmel

Himme!

JUN & 3 1983

50.9

LARC TECH LIBRARY

2725 488 408

02-02-1884 11:58

If you have any questions concerning this matter, please call Mr. William L. Bimkins at extension 3281.

Jess G. Ross 2898

> distribution: SDL 031

acı

NASA Scientific and Technical Information Facility P.O. Box 8757 EWI Airport, MD 21240

NASA--NIS-5/Security 180a/RIAD 139a/TULAO

139A/HLSImkinsielf 06/15/83 (3281)

139NJS D 6-15: 13

at-01 HEADS OF DRGANIZATIONS
HESS. JAME S..
BEG SITS JIAM

804 884 5375

1



NATIONAL ADVISORY COMMITTEE FOR AERONAUTICS

RESEARCH MEMORANDUM

PROPERTIES OF LOW-CARBON N-155 ALLOY BAR STOCK

FROM 1200° TO 1800° F

By J. W. Freeman and A. E. White

SUMMARY

The data in this report are the results of an investigation initially undertaken to establish rupture and total-deformation strengths ("design" data) at 1200° to 1800° F for typical commercial treatments of low-carbon N-155 alloy. Rupture data are reported for bar stock from two heats at 1200° , 1350° , and 1500° F and for one of the heats at 1650° and 1800° F in several conditions of prior treatment. Data on creep and on stress and time for total deformation are partially complete at 1200° , 1350° , and 1500° F.

The results show that there were large differences in strength between the heats of bar stock at temperatures above 1200° F except when a 2200° F solution treatment was used. This was due to steeper stress - rupture-time curves and lower creep resistance of one of the heats, except when they were solution-treated at 2200° F. This caused wide variation in the temperatures at which one type of treatment became superior to another. Likewise the comparison between treatments was complicated by the relationship changing depending on the criterion of strength used. Because the variation between heats indicated that the data would be of questionable value as typical design data, the work was stopped with the data on stress and time for total deformation somewhat incomplete and with very little duplication between the two heats.

The influence of prior treatment on the rupture data and design data, however, was in accordance with expectations. That is, hot-coldwork was beneficial up to a limiting temperature depending on the time period and criterion of strength used. This beneficial effect of hot-cold-work was maintained to temperatures as high as 1600° F for rupture in 1000 hours. Hot-rolled material was variable and intermediate to low in strength. Solution-treating and aging produced the highest strengths at the more elevated temperatures. The difference in properties of the two heats, however, caused wide differences in the transition temperatures of superiority of type of treatment.

The heat with the lower rupture strengths at 1350° and 1500° F developed a sigma-type phase as separate grains during testing. The reason why the sigma-type phase should be so detrimental or why it should form in one heat and not in the other is not known. There was no difference in chemical composition to account for the formation of a sigma-type phase. The only difference noted was a 1725° F finishing temperature during hot-rolling as compared with a 1910° F finishing temperature for the stock which did not develop sigma-type-phase grains. This investigation therefore indicates that hot-working conditions prior to final treatments have a pronounced effect on properties unless the final heat treatment involves a solution treatment at high temperatures to minimize variations in hot-working conditions.

INTRODUCTION

The development and use of forged heat-resisting alloys for turbojet engines have been complicated by several metallurgical factors.
Properties of such alloys at elevated temperatures vary depending upon
the type of final treatment. As-hot-worked, aged, hot-cold-worked,
solution-treated, solution-treated and aged, or solution-treated and
hot-cold-worked conditions are all used in practice. The relative—
properties for the various types of final treatment are changed considerably by the temperatures and time periods of heat treatment, the
temperatures and amount of hot-cold-work, and the treatment prior to
hot-cold-work. The situation is further complicated because the
relative properties vary considerably for the various conditions of
prior treatment depending on the temperature, time period, and criterion
of strength considered in test evaluation of the alloys or in service.

As a result, existing data for forged heat-resisting alloys are often contradictory. It is difficult to define the influence of chemical composition and practically no data exist which show consistent variations in properties for systematic changes in chemical composition. The best treatments for various service conditions are likewise not well-defined. In practical application of the alloys, strength at high temperatures is often substantially less than that predicted by test data and frequently is quite variable.

Previously an extensive study was made of the influence of type and conditions of prior treatment on the rupture properties of one heat of low-carbon N-155 alloy at 1200° F. (See reference 1.) The investigation covered by the present report initially had two objectives. One was to extend the previous investigation to include temperatures up to 1800° F. Complete evaluation of all the variations in conditions of prior treatments would have required too many tests to be practical.

Consequently it was decided to limit the testing to those conditions of prior treatment which seemed most usable in service on the basis of the results at 1200° F. Concurrently the NACA initiated a program to obtain design data in the form of curves of stress against time for total deformation at 1200°, 1350°, and 1500° F for common industrial treatments of the alloy. Again, in order to restrict the testing program to a manageable size, effort was concentrated to establish the curves for total deformations of 0.2 and 1 percent.

The available bar stock from the heat used for the tests of reference 1 was not sufficient for the entire program and it was necessary to procure more from a new heat. Duplicate rupture tests were carried out in part on both heats in order to include the two different initial materials as a variable.

The rupture tests on the second heat quickly demonstrated that there were considerable differences in the properties of the two heats except when a solution temperature of about 2200° F was used prior to testing. Apparently differences in the hot-working conditions prior to final treatment had a pronounced influence on the properties except after a high-temperature solution treatment. Recognition of this variable indicated that the design curves for the "typical" conditions of treatment would have only limited value. It also emphasized the need for concentration of effort on the fundamental crystalline structures which control properties at elevated temperatures. For this reason the testing for establishing design curves was stopped with many of the curves not too well-established.

This report presents the data obtained in the investigation with a minimum of fundamental interpretation. The main value of the results is the demonstration of the marked differences in properties at various temperatures, time periods, and criterions of strengths between two lots of stock unless they are heat-treated at high temperatures to minimize initial variations in the original test stock.

The investigation was conducted by the Engineering Research Institute of the University of Michigan under the sponsorship and with the financial assistance of the National Advisory Committee for Aeronautics.

TEST MATERIAL

Low-carbon N-155 alloy bar stock from two commercial heats was used in this investigation. Chemical analyses of the two heats were as follows:

Heat	Analysis by -				Chemi		omposite	tion			
	(1)	С	Mn	Si	Cr	Ni	Co	Мо	W	Ср	N
30276	U.C. U.M.	0.12	1.64 1.64	0.39 	21.33 21.18	18.88 18.59	18.60 18.99	3.21 3.14	1.97 2.05	1.10 1.12	0.12
A1726	U.C. U.M.		1.63 1.43		21.22 20.73						

1U.C. - Universal Cyclops Steel Corporation.

U.M. - University of Michigan.

The hot-working procedures used in the fabrication of the bar stock were reported by Universal Cyclops to have been as follows:

Heat 30276: The approximately 600-pound (8- by 7-inch) ingot was hammer cogged to 2-inch-square billets with 2050° F as maximum and 1750° F as minimum working temperatures. The billets were hot-rolled to 7/8-inch-square bars in one heat from 2075° F with a finishing temperature of 1725° F.

Heat A1726: The 13-inch-square ingot was hammer cogged in 15 heats to a 2-inch-square billet with 2070° F as maximum and 1730° F as minimum working temperature. The billet was hot-rolled to 7/8-inch-broken-corner-square bars in one heat from 2110° F with a finishing temperature of 1910° F.

EXPERIMENTAL PROCEDURE

The bases for selecting the prior treatments to be studied were:

- 1. Types of treatment widely used in practice and therefore of greatest interest, or outstanding properties based on reference 1.
- 2. Selection of conditions of treatment were based on common practice or upon the optimum conditions indicated by the data in reference 1.

The treatments used, outlined in detail in table I, included:

l. As-rolled, as-rolled and aged, and as-rolled and hot-coldworked because they represent common commercial practice. Other work (see reference 2) at temperatures above 1200° F had indicated that an age at 1350° F for 50 hours would be the most acceptable aging treatment. Reductions of 15 percent at 1200° F are fairly representative of commercial hot-working conditions. This hot-cold-worked condition was also included because as-rolled and hot-cold-worked material generally has adequate elongation in the rupture test at 1200° F as well as high strength.

- 2. Solution-treated and solution-treated plus aged materials because they are common treatments and because they were expected to have superior properties at the more elevated temperatures. Generally the higher the temperature of solution treatment the better is the strength of temperatures above 1350° F. The work of reference 1 however had indicated that for unaged material the maximum temperature of solution-treating was 2100° F without abnormally low elongation in the rupture test at 1200° F. Subsequent aging, however, improves elongation, the work of reference 1 indicating that treatment at 2200° F followed by aging at 1400° F for 24 hours gave the best combination of strength and ductility. Establishment of the design data at 1200° F on material aged for 50 hours at 1350° F was undertaken early in the program to conform with the treatments being used at other laboratories for testing at temperatures above 1200° F. (See reference 2.)
- 3. Solution-treated and hot-cold-worked material because this type of treatment gives by far the highest rupture strengths at temperatures on the lower end of the range being considered. The solution temperature of 2050° F was selected on the basis of the work in reference 1 which showed that 2050° F was the maximum solution temperature prior to hot-cold-work without excessive brittleness in the rupture test. A reduction of 15 percent at 1200° F was used for the reasons mentioned previously. The reduction of 25 percent was included to show the influence of increased reduction.

The room-temperature physical properties resulting from these treatments are included as table II as a matter of interest to those using the materials.

Data on stress and time for total deformation (elastic plus plastic deformation) were originally to be obtained at 1200°, 1350°, and 1500° F for all nine treatments. Sufficient tests were to be made so that curves of stress against time for rupture and for 0.2- and 1.0-percent total deformation could be plotted out to 2000 hours. Because of the excessive testing involved the program was reduced to include only

as-rolled, as-rolled plus 15-percent hot-cold-worked, 2100° F water-quenched, 2200° F water-quenched plus aged 24 hours at 1400° F, and 2050° F water-quenched plus 15-percent hot-cold-worked.

Total-deformation data were not duplicated between the two heats except at 1200° F. The evident wide difference in rupture properties between the two heats except after a 2200° F solution treatment indicated that duplicate data would have only limited utility. Furthermore, the testing program was terminated before most of the design curves were well-established.

Rupture testing was quite complete. Duplicate tests, however, between the two heats were limited to the five treatments finally selected for extensive testing. Rupture tests at 1650° and 1800° were also limited to one heat. Creep strengths reported were based on tests made to obtain design data.

Solution treatments were carried out in a gas-fired furnace. Aging was done in laboratory electric resistance furnaces. Hot-cold-working was accomplished by rolling 8-inch-long bars in a 5-inch, two high rolling mill.

Rupture tests were run in individual stationary units applying the stress through a simple-beam and knife-edge system. Rupture test specimens were 0.250 inch in diameter with a gage section 1 inch in length in all cases except the 1200° F tests on heat 30276 which were made on 0.160-inch-diameter specimens. The latter test data were taken from the previous work (reference 1). Time-elongation data were obtained during the rupture tests both by the drop-of-the-beam method and, in cases where design data were being obtained from rupture tests, by means of modified Martens' type extensometers with a sensitivity of 0.00005 inch per inch.

Creep tests were conducted on 0.505-inch-diameter specimens with a 2-inch gage length in units similar to those used for rupture testing. Duration of the creep tests varied up to 1700 hours depending on the total-deformation data needed from the particular test. The creep data were obtained by means of modified Martens' type extensometers with a sensitivity of 0.000003 inch per inch. In all cases the total deformation reported included the elastic deformation when the load was applied as well as the subsequent plastic deformation.

Room-temperature hardness, tensile, and impact tests were obtained before and after creep testing of material from heat A1726 as a measure of the structural stability of the alloy during creep testing.

Metallographic samples of the original material and the completed test specimens were prepared for observation and photomicrographs were taken of representative samples.

The indication of a sigma-type-phase formation during rupture testing was checked by X-ray diffraction methods. The procedure used was the low-angle technique outlined by Barnett and Troiano (reference 3). Rods approximately 1/16 to 1/32 inch in diameter were ground from two specimens after rupture testing. After electrolytic etching in ferric chloride to expose the sigma-phase particles, X-ray exposures of chromium Ka radiation up to 24 hours' duration were made using a Debye-Scherrer-Hull camera. The diffraction lines obtained were compared with those reported in references 4 and 5.

The data obtained are presented as a series of tables and figures which show rupture, stress and time for total deformation, creep, and stability characteristics.

Rupture Test Data

The data obtained from the rupture tests are given in tables III to VII. Curves of logarithmic stress against logarithmic rupture time are shown as figures 1, 2(a), 2(b), and 2(c). The rupture strengths of table VIII were read or extrapolated from these curves. The influence of testing temperature on the rupture strength and elongation is shown by figures 3 and 4.

The relationships between properties in the rupture test and prior treatment as influenced by testing temperature were complicated. The relative properties varied depending on the time period for rupture. The two heats also had different levels of properties because the slopes of their stress - rupture-time curves were different. All of the data have been compared in figures 5, 6, 7, and 8 with those for the samples solution-treated at 2200° F and aged for 24 hours at 1400° F on the assumption that this treatment minimized differences due to initial variations in the original hot-rolled stocks.

The variation in rupture strength between the two heats was small after solution-treating at 2200° F and aging at 1400° F for 24 hours. Furthermore reference to figure 2 shows that the slopes of the stress - rupture-time curves were similar. It appeared that solution-treating at increasing temperatures from 2050° to 2200° F reduced the difference in slope between the stress - rupture-time curves which existed for the stock from the two heats in the as-rolled condition. The 2200° F solution treatment did not, however, prevent wide differences in elongation in the rupture test. In spite of the difference in elongation it appears

that the assumption of minimized variations by a 2200° F solution treatment and 1400° F age was reasonably good.

The relationship between prior treatment and testing technique was in accord with general experience. That is:

- 1. Above a limiting temperature the solution-treated or solution-treated and aged condition will have higher rupture strength than the hot-cold-worked condition. The temperature and degree of superiority of the hot-cold-worked condition decrease with the time period for rupture considered.
- 2. Solution-treated and hot-cold-worked materials have substantially higher rupture strengths for 100 and 1000 hours than solution-treated or solution-treated and aged materials up to temperatures ranging from 1450° to 1680° F. Increasing the percentage of hot-coldwork from 15 to 25 percent in the case of heat 30276 resulted in poorer maintenance of strength with both increasing time and temperature of testing.
- 3. The strengths of stock hot-cold-worked in the as-rolled condition are not maintained as well as those solution-treated prior to hot-cold-work when testing time or temperature is increased.
- 4. Strengths in the hot-worked condition also tend to fall off rapidly in comparison with those of solution-treated stock with increasing time or temperature for rupture and are particularly low at temperatures above 1350° to 1500° F.
- 5. General experience indicates that comparative rupture strength increases with solution temperature as the test temperature or time is increased. This effect was only minor in the case of heat 30276. The data suggest it would have been even less for heat Al726.
- 6. The influence of aging after solution treatment on the rupture strength at varying temperatures is not well-defined by the data. At 1200° F aging for 24 hours raised the level of the stress rupture-time curves. There was practically no effect at 1350° F. At 1500° and 1650° F the short-time strengths were increased somewhat by aging and the long-time strengths lowered because of steeper stress rupture-time curves after aging. At 1800° F there was no significant effect.

Aging of hot-worked material seems too complicated to define in view of the differences in hot-worked materials. In the case of heat 30276 aging at 1350° F for 24 hours had little effect at 1200° and 1500° F while it reduced the slope of the stress - rupture-time curve at 1350° F. Considerably different effects would probably have been obtained by aging the as-rolled stock from heat A1726.

7. The elongation data from the rupture tests showed that abnormally low elongation only results during testing at 1200° F. While differences exist at higher temperatures all elongations were at least 5 percent. Solution-treating or hot-cold-working resulted in elongation below 5 percent at 1200° F. The samples without hot-cold-work tended to reach a maximum in elongation at 1350° to 1500° F, whereas the hot-cold-worked materials tended to have increased elongation with increasing temperature of test. Consequently the hot-cold-worked conditions had higher elongation at the higher temperatures.

When the investigation was undertaken it was hoped that the temperatures of superiority of rupture strength for the various treatments would be defined. The results, however, show the optimum properties at intermediate temperatures vary depending on the response of the particular lot of bar stock to the treatment as well as the time period on which comparisons are based. At 1200° F the best properties were obtained by hot-cold-work in all cases. Solution-treated or solution-treated and aged was best at 1800° F. The temperature at which the change-over occurred varied widely for the two heats and the various prior treatments. The temperatures at which the material solution-treated at 2200° F and aged at 1400° F became superior to other conditions have been compiled in table IX to illustrate this finding.

Characteristics of Stress and Time for Total Deformation

Design curves showing the relationship between stress and time for various total deformations are shown in figures 9 to 15. The data for these curves were taken from time-elongation curves from the individual tests, as summarized in tables X to XIII.

Stresses for total deformations of 0.1-, 0.2-, 0.5-, and 1.0-percent total deformation and the transition to third-stage creep in 10, 100, 1000, and 2000 hours as defined by the design curves are summarized in tables XIV and XV. In a number of cases the data were not sufficiently complete at the time testing was terminated to define all the deformation strengths. The influence of prior treatment on the various deformation strengths at the three temperatures at which heat A1726 was tested is shown graphically by figure 16. The curves of stress and time for total deformation at 0.2- and 1.0-percent deformation for the various treatments are compared in figures 17 and 18.

The results of the total-deformation tests indicate that:

l. Hot-cold-work substantially increases the permissible stress for limited total deformations for time periods up to at least 2000 hours at 1200° and 1350° F but not at 1500° F.

- 2. As in the rupture tests the as-rolled stock from heat A1726 compared favorably with the solution-treated and the solution-treated plus aged conditions. In this condition heat 30276 was inferior at 1200° F, the only temperature at which it was tested.
- 3. There was only a small difference between material solution-treated at 2200° F and material solution-treated at 2200° F and aged for 24 hours at 1400° F.
- 4. Total-deformation data are a complex combination of initial deformation when the load is applied, the amount of first-stage creep, and the inherent creep resistance. The first two are more influential at the lower temperatures and higher total deformations considered. Differences due to prior treatment tend to become less as the amount of total deformation decreases. Creep resistance would have more influence at longer time periods than those considered in the investigation. Two cases where low creep resistance probably contributed to low comparative total-deformation strength were the material from heat 30276 in the hot-rolled condition and in the solution-treated plus hot-cold-worked 25-percent condition.
- 5. The wide difference in the two hot-rolled materials at 1200° F probably indicates that there would have been wide differences in the total-deformation characteristics if the two had been tested with other treatments. As in the rupture test this difference probably would have been a minimum after a 2200° F solution treatment. Except for the solution-treated and aged stock there could be considerable variation in deformation strengths from those in this report. The results presented, however, are probably qualitatively correct insofar as the trends from the various treatments are concerned.

Creep Characteristics

The creep rates measured from the total-deformation tests are summarized in tables XVI and XVII. In most low-stress tests the creep rates were continuing to decrease with increased time of testing. These tables also show the minimum creep rates and time of transition to third-stage creep for the rupture tests.

The curves of logarithmic stress against logarithmic creep rate for heat 30276 are shown in figure 19. The hot-cold-worked condition had the highest creep resistance, the solution-treated and aged was intermediate, and the hot-rolled was weakest. The stress - creep-rate data were particularly erratic for the hot-cold-worked samples.

In figures 20(a) and 20(b), the stress-creep data at 1200°, 1350°, and 1500° F are assembled for the four conditions of treatment of heat A1726. These curves also show the pronounced influence of testing time on the observed creep rates. The pronounced influence of time on the observed creep rates leads to complex relations of creep strength, temperature, and treatment. It is also probably a contributing factor to the apparent erratic nature of the stress - creep-rate data. The stress - creep-rate characteristics of the hot-cold-worked material tested were more erratic than the other treatments particularly at 1500° F. Either the bars were not uniformly hot-cold-worked or, more probably, the instability of hot-cold-worked material at higher temperatures resulted in variability.

In general the number of tests run on heat A1726 was not sufficient to establish creep strength for 0.0000l percent per hour. The strengths for 0.000l percent per hour were, however, quite well-established, except in those cases where creep rates were changing rapidly with the testing time. The creep strengths indicated by the available data are summarized in table XVIII. The influence of temperature and treatment on the creep strengths, shown graphically in figure 16, was:

- l. Relative creep strengths were not well-defined at 1200° F, principally because the tests were incomplete for solution-treated and for solution-treated and aged material. The indications were, however, that 15-percent hot-cold-work at 1200° F nearly doubled the creep strengths. There was a wide difference in the creep resistance of the two heats in the as-rolled condition.
- 2. At 1350° F the superiority of the hot-cold-worked condition was greatly reduced for a rate of 0.000l percent per hour and had disappeared for a rate of 0.000l percent per hour. There was little difference between solution-treated (2100° F), solution-treated (2200° F) plus aged, and the hot-rolled stock for a rate of 0.000l percent per hour. The hot-rolled was inferior at 0.0000l percent per hour.
- 3. At 1500° F the data indicate that hot-cold-work was of no benefit for a rate of 0.0001 percent per hour and was detrimental at slower creep rates. Aging appeared to be detrimental to creep strength at this temperature. As-rolled material was decidedly inferior.

The creep strengths for a rate of 0.0001 percent per hour are compared in table XIX with the extrapolated rupture strengths for fracture in 10,000 hours. This creep strength is often extrapolated as the stress to cause 1-percent total deformation in 10,000 hours. The rupture data in table XIX show that the creep strengths approached the rupture strengths for the hot-cold-worked materials at all three temperatures. The difference was rather small at 13500 F for the

solution-treated and for the solution-treated plus aged conditions. The creep and rupture strengths were equal for all four conditions tested at 1500° F. When the extrapolated rupture strengths approach the creep strength, there is considerable doubt regarding the reliability of extrapolation of the creep strength. Probably third-stage creep will occur prior to 10,000 hours with a consequent greater elongation in 10,000 hours than 1 percent, or the materials will fracture after about 1-percent creep.

Stability Characteristics

Tensile, impact, and hardness tests and metallographic examination at room temperature were used to investigate structural stability during testing. The physical tests were confined to heat A1726 because unfractured specimens were available from all three temperatures of testing for total deformation.

The results of the tensile, impact, and hardness tests are in table XX. In all conditions of treatment there was a marked and progressive drop in ductility and impact strength after creep testing at 1200° and 1350° F. After testing at 1500° F these properties were either slightly less or slightly higher than after testing at 1350° F. It is particularly interesting to note that the solution-treated and hot-cold-worked stock underwent no greater change in this respect than the solution-treated and aged. The as-rolled material had slightly better ductility after testing than the other conditions.

The strength and hardness of all conditions, except the hot-cold-worked, were increased by creep testing at 1200° and still further increased by testing at 1350° F. Testing at 1500° F did not produce as much increase in strength and hardness. The hot-cold-worked stock progressively decreased in strength and hardness with increasing testing temperatures.

These fairly pronounced changes in strength and ductility as a result of testing indicate that the alloy undergoes considerable structural alteration during exposure to temperature and stress. These apparently are much greater at 1350° and 1500° F than at 1200° F for time periods up to 2000 hours. Aging for 24 hours at 1400° F after solution-treating at 2200° F apparently did very little to stabilize the structure. The hot-cold-worked material apparently underwent considerable relief of strain hardening during testing.

A metallographic examination was made of all of the original materials, of the longest duration rupture specimens, and of representative unfractured specimens from the design-data tests. The structural

changes were of a different type in heat 30276 from those in heat A1726 as is shown by the representative photomicrographs of figures 21 to 28. Both materials appeared to undergo both general and intergranular precipitation during testing at 1200° F. At 1350° and 1500° F, however, heat 30276 showed the presence of an agglomerated constitutent which microscopically had all the characteristics of sigma phase. The precipitation in heat A1726 remained fine and well-dispersed after testing at these temperatures, the size of the particles merely increasing with testing temperatures.

The two rupture specimens from heat 30276 showing the strongest indication of sigma phase in the metallographic examination were checked by X-ray diffraction for sigma phase with the results shown in table XXI. A considerable number of lines corresponding closely in spacing to published diffraction values for sigma phase were found for the specimen tested at 1500° F and four lines for the specimen tested at 1350° F. These results confirm the presence of a sigma-type phase indicated by the metallographic examination. Because a number of the elements present in low-carbon N-155 alloy have been shown to form the sigma-phase structure (reference 4) and because the composition of the phase was not determined, the term "sigma-type phase" is used in this report. In addition to the sigma-phase lines, the columbium carbide in the low-carbon N-155 samples also gave diffraction lines.

The amount of sigma-type phase which formed in the samples from heat 30276 decreased with increasing solution temperatures and the structures after a 2200° F solution treatment were similar to those from heat A1726. Hot-cold-work apparently increased the tendency for the sigma-type phase to form in heat 30276. There were differences in grain size between the two heats although this factor seemed small in comparison with the differences in precipitation characteristics.

DISCUSSION OF RESULTS

This investigation was originally undertaken for the express purpose of providing representative design data for several commercial treatments. The results show that there can be quite wide differences in properties between two lots of the alloy which appear to be the same when compared on the basis of chemical composition or the usual room temperature physical properties.

The results do show that the usual generalizations regarding the effect of prior treatment on properties at high temperatures are correct. There can, however, be wide differences in the actual magnitude of the rupture, total deformation, or creep strengths as well as in ultimate elongation to fracture for different lots of the alloy.

Furthermore there can be wide variations in the temperature at which one type of treatment provides superior strength to another type. In this investigation, for instance, hot-cold-work provided superior rupture strength in one heat even for prolonged time periods at temperatures above 1500° F, while the other heat was inferior to solution-treated and aged material at temperatures below 1500° F at fairly short time periods. The as-rolled materials were particularly divergent in this respect. Thus there is not a single limiting temperature, even for a specific measure of strength, of superiority of one type of treatment over another for various lots of test stock.

The lower strength of heat 30276 than of heat A1726 at the higher temperatures and longer time periods was associated with the formation of a sigma-type phase during testing. The conditions under which it formed, the influence of prior treatment, and the effect on properties at high temperatures were similar to those observed in 18-8+Cb and 25-20 steel. (See reference 6.) The phase forms most extensively at about 1350° F, forms only slightly at 1200° F, and forms larger grains at 1500° F. High stresses and deformation, as in rupture testing, promote its formation. Increasingly higher solution temperatures reduce the amount of the phase which forms and finally eliminate it, at least as separate grains. Cold-work accelerates its formation. The formation of the sigma-type phase as separate grains is associated with steep stress - rupture-time curves, lowered creep resistance, and increased elongation in the rupture test.

Apparently solution-treating at 2200° F was nearly high enough to prevent grains of sigma-type phase from forming in heat 30276. This then could account for the similarity in rupture properties to those of heat A1726 after this treatment. Some sigma-type phase formed, however, causing some loss in creep strength which was offset by the increased elongation so that the rupture strengths were about equal after the 2200° F treatment.

The absence of identifiable grains of the sigma-type phase after testing at 1200° F indicates that either it does not form at that temperature or the rate of formation is extremely slow. The strength of heat 30276 was therefore comparable with that of heat A1726 at 1200° F. The reason why formation of sigma-type phase at higher temperatures apparently reduces strength in the rupture and creep tests is not evident from available information. The accumulation of precipitants into the grains of sigma phase leaving clean grain boundaries could account for the increased elongation.

The reason for the formation of sigma-type phase in specimens from heat 30276 and not in those from heat A1726 is not apparent. There is nothing in the chemical composition to account for the difference. The

only difference in the reported history of fabrication of the two was a finishing temperature of 1725° F for heat 30276 and 1910° F for heat A1726. It is therefore suggested that differences in hot-working conditions were responsible for the differences in properties of the two heats. It appeared as if a 2200° F solution treatment minimized such differences. It is believed, however, that the 2200° F treatments should not be relied upon for this purpose pending further verification over a wider range of hot-working conditions.

CONCLUSIONS

Rupture data are reported at 1200°, 1350°, and 1500° F for two heats of low-carbon N-155 alloy bar stock in several typical conditions of treatment. The rupture test data were extended to 1650° and 1800° F for one of the heats. Design data in the form of the relationship between stress and time for various amounts of total deformation up to 1 percent were partially obtained for various typical treatments at 1200°, 1350°, and 1500° F. The results showed that:

- l. The two heats differed in strength except when they were both solution-treated at 2200° F and aged at 1400° F prior to testing. The differences were small at 1200° F but pronounced at higher temperatures.
- 2. One heat had much steeper stress rupture-time curves than the other. This was associated with the formation of grains of a sigma-type phase in the microstructure during testing at 1350° and 1500° F. The only difference observed between the two heats to which this difference in behavior could be attributed was a lower finishing temperature during the hot-rolling of the bar stock.
- 3. The results in general confirmed previous experience in that hot-cold-worked material loses its superiority in strength above some limiting temperature. This temperature decreases with the time period considered. The superiority also becomes less as the total deformation on which comparisons are based decreases. The hot-cold-worked condition, however, was found to maintain superiority to much higher temperature than had been thought to be the case. As-hot-rolled materials can have very variable properties, apparently associated with variations in hot-working conditions, little difference in solution-treated or solution-treated and aged material was observed.
- 4. While the general relations between prior treatment and properties at various temperatures are known the actual magnitude of the properties between different lots of an alloy can vary widely.

Consequently it is not possible to establish "typical" properties for various treatments except after high-temperature solution treatments which minimize differences in prior hot-working conditions.

Engineering Research Institute
University of Michigan
Ann Arbor, Mich., May 31, 1950

REFERENCES

- 1. Freeman, J. W., Reynolds, E. E., Frey, D. N., and White, A. E.:
 A Study of Effects of Heat Treatment and Hot-Cold-Work on
 Properties of Low-Carbon N-155 Alloy. NACA TN 1867, 1949.
- 2. Cross, Howard C., and Simmons, Ward F.: Heat-Resisting Metals for Gas-Turbine Parts. Symposium on Materials for Gas Turbines, A.S.T.M., 1946, pp. 3-51.
- 3. Barnett, W. J., and Troiano, A. R.: X-Ray Identification of Sigma Phase in 25-20 Cr-Ni Stainless. Metal Progress, vol. 53, no. 3, March 1948, pp. 366-367.
- 4. Duwez, P., and Baen, S. R.: X-Ray Study of the Sigma Phase in Various Alloy Systems. Preprint 47, A.S.T.M. 1950.
- 5. Anon.: X-Ray Diffraction Data Cards. A.S.T.M. (Philadelphia).
- 6. Clark, C. L., and Freeman, J. W.: The Apparent Influence of Grain Size on the High Temperature Properties of Austenitic Steels. Trans. Am. Soc. Metals, vol. 38, 1947, pp. 148-179.

TABLE I.- HEAT TREATMENTS AND TESTING CONDITIONS USED TO OBTAIN RUPTURE AND DESIGN DATA

	Tests con	iducted and temper	ratures of	testing	
Treatment	Rupture	e tests	Design-data tests		
	Heat 30276	Heat Al726	Heat 30276	Heat A1726	
As-rolled .	1200° to 1500° F	1200° to 1800° F	1200° F	1200° to 1500° F	
As-rolled; aged at 1350° F for 24 hr	1200° to 1500° F				
As-rolled; 15% hot-cold-work at 1200° F	1200° to 1500° F	1200° to 1500° F			
2100° F, 1 hr, W.Q. ²	1200° to 1500° F	1200° to 1800° F		1200° to 1500° F	
2200° F, 1 hr, W.Q.	1200° to 1500° F		~~~~~		
2200° F, 1 hr, W.Q.; aged at 1400° F for 24 hr	1200° to 1500° F	1200° to 1800° F	~~~~~	1200 ⁰ to 1500 ⁰ F	
2200° F, 1 hr, W.Q.; aged at 1350° F for 50 hr	1200° F		1200° F		
2050° F, 2 hr, W.Q.; 15% hot-cold-work at 1200° F	1200° to 1500° F	1200° to 1800° F	~~~~~	1200 ⁰ to 1500 ⁰ F	
2050° F, 2 hr, W.Q.; 25% hot-cold-work at 1200° F	1200° to 1500° F		1200 ⁰ F		

 $^{1}\mathrm{Test}$ temperatures were 1200°, 1350°, 1500°, 1650°, and 1800° F; table shows range of temperatures at which tests were made.

²W.Q. - Water-quenched.

TABLE II. - ROOM-TEMPERATURE PHYSICAL PROPERTIES OF LOW-CARBON

N-155 ALLOY BAR STOCK FROM TWO HEATS

Heat	Heat Treatment beriness at:	Tensile atrength	Of£1	set yield strem (psi)	ngth	Proportional limit	Elongation in 2 in.	Reduction of area	
	(1)	nardness	(psi)	0.02 percent	0.1 percent	0.2 percent	(psi)	(percent)	(percent)
30276 A1726	As-rolled As-rolled	233 228	128,500 127,200	72,500 63,000	76,500 73,200	78,500 77,400	57,500 46,500	40.5 40	55.7 51.0
30276	AR+Ag	214	124,500	52,700	62,500	66,300	37,500	36	38.2
30276 A1726	AR+HCW AR+HCW	328 337	156,700	121,500	135,500	140,000	82,500	22,	43.3
30276 A1726	ST(2100) ST(2100)	197 207	117,250 117,000	38,500 45,700	52,000 52,800	57,000 56,300	17,500 36,000	47 50	63.2 67.6
30276	ST(2200)	205	115,750	42,000	53,000	57,000	25,000	46.5	64.3
30276 A1726	ST+Ag(1400) ST+Ag(1400)	221 228	119,250 118,900	48,000 47,800	61,000 55,800	<i>6</i> 5,000 59,400	25,000 3.7,000	32 38	39.8 46.8
30276	ST+Ag(1350)	213	118,000	50,000	57,500	60,000	32,500	36	41.9
30276 A1726	ST+15%HCW ST+15%HCW	296 321	145,500 144,000	110,500 107,500	120,500 117,200	124,500 120,300	65,000 97,500	26 24	53.2 57.0
30276	817+2596HCW	318	156,000	104,000	130,000	137,500	70,000	22	48.1

l_{Treatments:}

TABLE III.- RUPTURE TEST DATA AT 1200° F FOR LOW-CARBON N-155 ALLOY BAR STOCK

Treatment	Heat	Stress (psi)	Rupture time (hr)	Elongation in 1 in. (percent)	Reduction of area (percent)
As-rolled	30276	855,000 850,000 845,000 840,000	8.5 75 252 610	^b 9.5 17 19 3 ⁴	8.5 23.3 23.3 48.3
As-rolled	A1726	50,000 49,000 45,000	53 80 472	c ₂ c ₅ 12	10.2 6.1 11.3
As-rolled + aged 24 hr at 1350° F	30276	⁸ 50,000 ⁸ 45,000 ⁸ 41,000	123 241 411	12 38 33	岁 .2 55.3 55.3
As-rolled + 15% hot-cold- work at 1200° F	30276	⁸ 60,000 ⁸ 55,000 ⁸ 50,000	157 325 845	6 8 6.5	20.6 24.1 17.9
As-rolled + 15% hot-cold- work at 1200° F	A1726	60,000 57,500 55,000	63 94 712	1 1 1	3.5 2.2 2.8
l hr, 2100° F, W.Q.d	30276	a50,000 a45,000 a40,000	35 141 1003	ъ9 ъ6 16	17.8 10.9 17.8
l hr, 2100° F, W.Q.	A1726	45,000 43,000 40,000 37,000	17 136 725 1845	^b 8 Ա Ե <u>Ա</u> 13	18.6 7.7 11.8 16.4
1 hr, 2200° F, W.Q.	30276	a50,000 a45,000 a40,000 a37,500	4 29 247 1500	b5 4 b6	26.7 15.6 6.2
l hr, 2200° F + 24 hr at 1400° F	30276	850,000 850,000 847,000 845,000	49 118 133 398	^b 12 14 18 21	18.3 17.8 23.3 30.8
l hr, 2200° F, W.Q.; 24 hr at 1400° F	A1726	50,000 47,000 46,000 44,000	66 101 144 446	10 9 10 10	11.7 8.8 9.8 14.5
l hr, 2200° F, W.Q.; 50 hr at 1350° F	30276	⁸ 50,000 ⁸ 45,000 ⁸ 40,000 35,000	53 145 479 3301	^b 12 ^b 10 ^b 14 19	13.3 16.7 21.2 21.6
2 hr, 2050° F, W.Q.; 15% hot- cold-work at 1200° F	30276	⁸ 60,000 ⁸ 55,000 ⁸ 52,000	167 389 1556	1 8 4	1.5 21.0 16.4
2 hr, 2050° F, W.Q.; 15% hot- cold-work at 1200° F	A1726	60,000 55,000 52,000 49,000 49,000 48,000	14.5 147 270 210 137 942	2 5 2 2 03 03 1.5	7-5 3-5 3-0 3-5 5-6 1.8
2 hr, 2050° F, W.Q.; 25% hot- cold-work at 1200° F	30276	⁸ 70,000 ⁸ 65,000 ⁸ 60,000 ⁸ 55,000	19 200 493 1142	2 4 5 5	5.7 11.0 18.9 21.6

80.160-in.-diam. test specimen. All other specimens were 0.250-in.-diam. bFractured in gage mark. cFractured in fillet. dW.Q. - Water-quenched.

TABLE IV.- RUPTURE TEST DATA AT 1350° F FOR LOW-CARBON N-155 ALLOY BAR STOCK

Treatment	Heat-	Stress (psi)	Rupture time (hr)	Elongstion in 1 in. (percent)	Reduction of area (percent)
As-rolled	30276	30,000 25,000 20,000	130 264 769	42 41 31	45.3 48.1 35.6
As-rolled	A1726	35,000 32,500 30,000	86 156 623	18 36 23	21.6 35.5 19.9
As-rolled + aged 24 hr at 1350° F	30276	30,000 22,000 20,000	51 351 639	50 38 30	62.5 50.2 36.8
As-rolled + 15% hot- cold-work at 1200° F	30276	35,000 30,000 22,000	95 172 499	19 19 12	27.1 22.6 14.6
As-rolled + 15% hot- cold-work at 1200° F	A1726	40,000 35,000 30,000 25,000	66 208 456 1693	6 8 5 5	6.9 9.9 4.1 3.2
1 hr, 2100° F, W.Q.ª	30276	30,000 25,000 22,000	117 432 997	35 21 23	38.4 25.9 32.5
1 hr, 2100° F, W.Q.	A1726	30,000 29,000 27,000 25,000	75 222 400 696	10 31 35 24	11.8 26.1 33.5 26.1
1 hr, 2200° F, W.Q.	30276	32,000 30,000 27,000 25,000	65 165 328 439	^b 13 36 35 b16	18.8 36.0 31.4 21.0
l hr, 2200 ⁰ F + 2 ¹ 4 hr at 1 ¹ 400 ⁰ F	30276	33,000 30,000 25,000	32 136 726	35 47 32	35.9 46.9 37.3
l hr, 2200° F, W.Q. + 24 hr at 1400° F	A1726 .	35,000 33,000 30,000 26,000	48 62 226 441	25 16 30 44	24.9 13.9 34.6 41.6
2 hr, 2050° F, W.Q. + 15% hot-cold-work at 1200° F	30276	39,000 35,000 30,000	79 250 593	^b 13 9 5	27.2 17.4 7.3
2 hr, 2050° F, W.Q. + 15% hot-cold-work at 1200° F	A1726	38,000 36,000 33,000	183 300 564	7 12 9	13.1 17.6 14.4
2 hr, 2050° F, W.Q.; 25% hot-cold-work at 1200° F	30276	39,000 33,000 30,000	132 319 446	7 4 6	11.5 5.5 8.2

aw.Q. - Water-quenched. bFractured in gage mark.

TABLE V.- RUPTURE TEST DATA AT 1500° F FOR LOW-CARBON N-155 ALLOY BAR STOCK

Treatment	Heat	Stress (psi)	Rupture time (hr)	Elongation in 1 in. (percent)	Reduction of area (percent)
As-rolled .	30276	20,000 15,000 12,000 9,000	16.5 64 194 430	26 48 33 28	60.6 44.0 35.1 28.6
As-rolled	A1726	15,000 13,500 13,000 12,000	130 458 433 747	25 23 22 17	25.2 24.0 19.8 20.0
As-rolled + aged 24 hr at 1350° F	302 ₇₆	17,500 13,000 9,000	27 102 424	46 40 27	53.2 42.3 28.9
As-rolled + 15% hot- cold-work at 1200° F	30276	22,000 16,000 9,500 7,500	1 ¹ 4 53 272 ¹ 93	35 31 22 44	37.3 30.6 24.6 29.1
As-rolled + 15% hot- cold-work at 1200° F	A1726	20,000 17,000 15,000 12,000	81 240 256 890	12 12 8 10	10.5 11.8 6.9 5.6
1 hr, 2100° F, W.Q.ª	30276	17,500 15,000 13,000	97 334 577	40 43 34	54.5 51.9 42.7
l hr, 2100° F, W.Q.	A1726	18,000 16,500 15,000	109 241 672	58 47 45	54.8 51.4 47.4
l hr, 2200° F, W.Q.	30276	20,000 17,500 15,000	71 255 615	60 40 36	52.7 49.3 40.8
1 hr, 2200° F + 24 hr at 1400° F	30276	19,500 17,500 14,000	160 . 239 . 1131	48 32 23	54.7 50.5 31.3
1 hr, 2200° F, W.Q. + 24 hr at 1400° F	A1726	20,000 18,000 16,000 14,600	131 260 760 1033	33 23 52 33	39-4 31-9 47-0 44-1
2 hr, 2050° F, W.Q. + 15% hot-cold-work at 1200° F	30276	20,000 17,500 15,000 13,000	136 303 486 421	16 ^b 5.5 9 12	17.7 3.4 5.4 9.9
2 hr, 2050° F, W.Q. + 15% hot-cold-work at 1200° F	A1726	24,000 20,000 18,000	104 321 739	14 10 6	24.0 14.5 6.9
2 hr, 2050° F, W.Q. + 25% hot-cold-work at 1200° F	30276	20,000 18,000 13,000	100 129 340	10 °6 10	11.4 7.8 10.6

aw.Q. - Water-quenched. bFractured in fillet. cFractured in gage mark.

NACA

TABLE VI.- RUPTURE TEST DATA AT 1650° F FOR LOW-CARBON N-155 ALLOY BAR STOCK

Treatment	Heat	Stress (psi)	Rupture time (hr)	Elongation in 1 in. (percent)	Reduction of area (percent)
As-rolled	A1726	7,000 5,500 5,000	148 334 1051	19 19 8	20.5 16.9 11.3
1 hr, 2200° F, W.Q. ¹	A1726	9,000 8,400 8,000	192 359 647	39 22 25	39.2 23.1 27.9
l hr, 2200° F + 24 hr at 1400° F	A1726	13,000 9,000 7,400	46 385 1384	39 22 17	43.4 23.3 19.9
2 hr, 2050° F, W.Q. + 15% hot-cold-work at 1200° F	A1726	14,000 11,000 9,000 7,000 6,200 6,000 5,500	65 165 301 676 1419 676 977	8 11 8 7 (2) 24 11	15.4 8.6 9.1 9.2 7.0 18.3 15.4

¹W.Q. - Water-quenched. ²Piece of specimen near fracture—lost.

NACA RM 51B05 . . 23

TABLE VII.- RUPTURE TEST DATA AT 1800° F FOR LOW-CARBON N-155 ALLOY BAR STOCK

Treatment	Heat	Stress (psi)	Rupture time (hr)	Elongation in 1 in. (percent)	Reduction of area (percent)
As-rolled	A1726	3000 2300 1800 1500	115 228 563 847	36 (1) 40 10	24.0 14.0 12.5 21.2
l hr, 2100° F, W.Q. ²	A1726	6600 5500 5000 4500 4500 3900 3500 2800	37.5 108 223 243 272 317 424 2128	31 12 20 13 14 21 18 12	28.0 13.2 22.1 10.0 14.5 19.6 17.5 14.5
l hr, 2200° F + 24 hr at 1400° F	A1726	7000 5000 3800 3300	43.5 188 485 959	32 9 10 8	26.1 9.3 7.5 9.0
2 hr, 2050° F, W.Q. + 15% hot-cold-work at 1200° F	A1726	6000 4500 3300 3000 2100 1850	22.5 51 175 271 474 1004	24 24 24 21 30 29	28.9 26.0 16.0 14.7 19.0 16.0

Priece of specimen near fracture lost. 2W.Q. - Water-quenched.

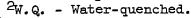




TABLE VIII.- COMPARATIVE HUPTURE PROPERTIES AT 1200° To 1800° F FROM TWO

HRATS OF LOW-CARBON N-155 ALLOY BAR STOCK

				Heat 30276	<u> </u>				Heat 41726		
Treatment	Temper- ature (°F)	Rupture strength (ps1)		Estimated rupture elongation (percent in 1 in.)		Rupture strength (psi)			Estimated rupture elongation (percent in 1 in.)		
		100 hr	1000 hr	10,000 ът	100 hr	1000 hr	100 per	1000 hr	10,000 hr	100 hr	1000 hr
As-rolled	1200 1350 1500 1650 1800	49,500 32,000 13,500	37,500 18,500 7,800	28,000 10,500 1,500	17 kg ko	34 31 28	48,000 34,000 15,500 7,800 3,150	43,000 29,000 11,500 4,100 1,400	38,000 24,000 8,600 2,100 620	5 20 25 19 36	12 23 17 · 8 10
2100° F, 1 hr, W.Q.1	1200 1350 1500 1650 1800	46,500 31,000 17,500	40,000 22,000 12,500	34,000 16,000 9,000	7 35 40	16 23 34	44,000 30,500 18,000 9,600 5,500	38,500 24,500 14,500 7,700 3,350	34,000 20,000 11,500 6,300 2,000	15 58 40 12	10 24 45 13
2200° F, 1 hr, W.Q.; 1400° F, 24 hr	1200 1350 1500 1650 1800	50,000 30,500 21,000	12,000 24,000 14,000	35,000 19,000 9,200	14 47 50	य ३२ २३	47,000 32,000 21,000 11,500 5,700	42,000 25,500 14,500 7,700 3,300	38,000 20,000 10,000 5,400 1,900	1.0 25 35 30 25	10 ht, 33 20 8
As-rolled; 15% hot-cold-work at 1200° F	1200 1350 1500	63,000 35,000 12,800	48,000 18,000 5,800	36,000 9,300 2,700	6 19 28	6 12 ¹ /4	59,000 37,500 18,700	54,000 27,000 11,800	50,000 19,000 7,300	1 6 12	1 5 10
2050° F, 2 hr, W.Q.; 15% hot-cold-work at 1200° F	1200 1350 1500 1650 1800	62,000 38,000 22,000	53,500 28,500 12,500	46,000 21,000 7,000	1 12 16	. 5 5 12	55,000 41,000 24,000 12,300 3,800	48,000 31,500 17,000 6,500 1,850	42,000 23,000 12,000 3,400 900	3 6 14 10 24	1.5 9 6 10 30
As-rolled; 1350° F, 24 hr	1200 1350 1500	51,000 27,000 13,000	35,500 18,500 7,400	24,000 13,000 4,300	12 45 41	33 30 27					
2200° 7, 1 hr, W.Q.	1200 1350 1500	42,000 30,500 19,000	36,000 23,500 14,500	34,000 18,000 11,000	4 35 50	6 16 36				-	
2050° F, 2 hr, W.Q.; 25% hot-cold-work at 1200° F	1200 1350 1500	66,000 41,500 20,000	56,000 25,000 9,000	15,000 15,000 3,900	3 7 10	5 6 10					
2200° F, 1 hr, W.Q.; 1350° F, 50 hr	1,200	46,500	39,500	32,000	п	15					

W.Q. - Water-quenched.



TABLE IX.- TEMPERATURES AT WHICH RUPTURE STRENGTHS OF OTHER TREATMENTS FELL BELOW THOSE RESULTING FROM WATER-QUENCHING FROM 2200° F AND AGING AT 1400° F FOR 24 HOURS

	Temperature (°F)									
Other treatments	100 hr fo	r rupture	1000 hr f	or rupture	10,000 hr for rupture					
	Heat A1726	Heat 30276	Heat A1726	Heat 30276	Heat Al726	Heat 30276				
2050° F, 2 hr, W.Q. + 15% hot-cold-work	1680	(>1500)	1600	1450	1560	1425				
2050° F, 2 hr, W.Q. + 25% hot-cold-work		1485		1365		1225				
As-rolled + 15% hot-cold-work	1455	1400	1400	1250	1360	1200				
As-rolled	1390	1375	1420	(1)	1460	(1)				
As-rolled + 24 hr at 1350° F		(1)		(1)	*** == ,= -	(1)				
2100° F, 1 hr, W.Q. ²	(1)	(1)	(3)	(1)	(4)	1200				
2200° F, 1 hr, W.Q.		(1)		(5)		(6)				

¹Strengths were less than those of the solution-treated and aged at all temperatures of testing.

²W.Q. - Water-quenched.

³Strengths were less than those of the solution-treated and aged up to 15000 F and the two were equal at higher temperatures.

Higher rupture strengths than those of the solution-treated and aged except at 1200° and 1800° F.

⁵Strengths were less than those of the solution-treated and aged up to 1375° F and the two were equal at higher temperatures.

⁶Lower rupture strengths than those of solution-treated and aged except at 1425° F and above.

Table x.- data on stress and time for total deformation at 1200° F for low-carbon N-155 allog bar stock from heat 30276

		Initial	Time (hr) for total deformation of -					nsition to -stage creep	Rupture data	
Treatment	Stress (psi)	deformation (percent)	0.1 percent	0.2 percent	0.5 percent	1.0 percent	Time (hr)	Deformation (percent)	Rupture time (hr)	Elonga- tion (percent)
As-rolled	10,500 11,000 12,500 13,500 15,000 18,000 20,000 22,500 25,000 30,000 40,000	0.044 .044 .048 .061 .066 .076 .100 .128 .128 .148 .331	990 361. 470 230 145 37	82450 81600 1310 432 90 41 28 9	2948 950 345, 236 110 3	B 1250 755 273 1 4	360 150	8.0 8.6	610 252	34 19
2200° F, 1 hr, W.Q.b + 50 hr at 1350° F	15,000 17,500 20,000 25,000 30,000 35,000 40,000	0.066 .074 .096 .123 .145 .221 .386	640 175 1,5	#2800 #175 73 -23 	*1045 190 26 2.5	620 110 13	2700 290	10.0 ት.95	3301 497	19 14
2050° F, 1 hr, W.Q. + 25% hot- cold-work at 1200° F	15,940 20,000 30,000 40,000 55,000 60,000 65,000	0.062 .092 .136 .169 ,232 .253 .274	a2500 10	82400 77 8 	44300 470 5 3-5	925 305 8	900 425 125	0.9 ⁴ 1.30 1.60	11½2 ½3 200	5 5 14

^aEstimated by creep data extrapolation.

bw.Q. - Water-quenched.

TABLE XI.- DATA OF STRESS AND TIME FOR TOTAL DEFORMATION AT 12000 F FOR LOW-CARBON H-155 ALLOY BAR STOCK FROM BEAT A1726

Treatment		Initial	Time (hr) for total deformation of -				Transition to third-stage creep		Rupture data	
	Stress (psl)	deformation (percent)	0.1 percent	0.2 percent	0.5 percent	1.0 percent	Time (hr)	Deformation (percent)	Rupture time (hr)	Klonga- tion (percent)
As-rolled	15,000 20,000 25,000 30,000 32,500 45,000 49,000 50,000	0.0550 .0905 .1162 .1273 .1432 .265 .313 .340	695	598 222 42 15	557 140	*2600 525 18 6 10	190	3.8	472 80 52.5	12 5 2
2100° 7, 1 hr, W.Q. ⁵	27,500 30,000 32,500 35,000 37,000 40,000	0.120 .160 .225 .320 .430 .870		72 14	540 370 265 68 2	*1680 1115 770 330 22 4	920	3.65	1845 725	13 4
2200° F, 1 hr, W.Q. + 24 hr at 1400° F	25,000 30,000 34,000 37,000 44,000 46,000	0.1125 .1366 .1759 .202 .636 .732		115 32 2.2 	1223 198 19 10	632 130 30 7 2	1150 265 94	1.6 4.5 5.4	446 144	10
2050° F, 2 hr W.Q. + 15% hot- cold-work at 1200° F	30,000 37,000 40,000 40,000 42,000 48,000 49,000 52,000 55,000	0.136 .168 .181 .181 .190 .217 .220 .220 .235		190 22 4 3 .5	C1315 C295 - C12 13 16 18 10 6	200 95 97	220	1.02	942 137 210 270 147	1.5 63 63 2.0 5.0

^aRstimated by creep data extrapolation. ^bW.Q. ~ Water-quenched. ^cO.25-percent deformation. ^dFractured in gage mark or fillet.

TABLE XII .- DATA ON STRESS AND TIME FOR TOTAL DEFORMATION AT 13500 F FOR LOW-CARBON N-155 ALLOY BAR STOCK FROM HEAT A1726

Treatment		Initial	Time (hr) for total deformation of -				Transition to third-stage creep		Rupture data	
	Stress (psi)	B deformation	0.1 percent	0.2	0.5 percent	1.0 percent	Time (hr)	Deformation (percent)	Rupture time (hr)	Elonga- tion (percent)
As-rolled .	10,000 15,000 20,000 22,500 30,000 32,500 35,000	0.0467 .0769 .0957 .1087 .1810 .240 .306	173 6 .5 	1555 225 25 5.4 	ठ १५८ १५८ ३१५६	1510 ⁸ 210 41 6 6	3 35 89	3.5 8.4	623 156 86	23 36 18
2100 ⁰ F, 1 hr, W.Q. ^b	12,500 15,000 17,500 20,000 25,000 27,000 29,000 39,000	0.053 .063 .074 .087 .095 .112 .135	19 8 1.2 5	192 47 8 12 1.6	464 67 66 6.5 9	725 273 20 13 6	405 220 95 65	7.4 9.0 6.4 5.7	696 460 222 75	24 35 31 10
2200° F, 1 hr, W.Q. + 24 hr at 1400° F	10,000 15,000 17,500 20,000 26,000 30,000	0.0483 .0731 .0781 .1024 .150	162 17 2,2	1240 186 60	650 68 7	273 14 5	270 99	7.2 6.2	հեյ 226	7/4 7/4
2050° F, 2 hr, W.Q. + 15% hot- cold-work at 1200° F	15,000 20,000 22,500 25,000 33,000 36,000 36,000	0.0680 .09to .1030 .1160 .149 .162 .175	75	1362 73 71 35 	700 24 17 6	*1850 80 52 24	1410 312 195 92	0.87 1.6 1.6 1.6	564 300 184	9 12 7

 $^{\rm a}_{\rm Estimated}$ by creep data extrapolation. $^{\rm b}_{\rm W}, \, q_{\star}$. Water-quenched.

TABLE XIII .- DATA ON STRESS AND TIME FOR TOTAL DEFORMATION AT 1500° F FOR LOW-CARBON N-155 ALLOY BAR STOCK FROM HEAT A1726

		Initial deformation (percent)	Time (hr) for total deformation of -					nsition to -stage creep	Rupture data	
Treatment	Stress (ps1)		0.1 percent	0.2 percent	0.5 percent	1.0 percent	Time (hr)	Deformation (percent)	Rupture time (hr)	Elonge- tion (percent)
As-rolled	7,500 9,000 10,000 10,500 12,000 13,000 13,500 15,000	0.0340 .0442 .0532 .0564 .0580 .0630 .0680	58 25 13 7 	550 197 75 	1520	18 8 	1225 400 110 260 72	0.42 3.4 2.6 3.5 6.6	747 432 458 130	20 22 23 25
2100° F, 1 hr, W.Q.a	8,500 10,000 12,000 15,000 16,500	0.0409 .0433 .0546 .181 .240	144 20 2.0	400 21 ¹ 4 33 	490 2	867 12 1	410 234 136	0.45 3.8 8.8	671 241	45 47
2200° F, 1 hr, W.Q. + 24 hr at 1400° F	7,000 9,000 10,000 12,000 14,600 16,000 18,000 29,000	0.0328 .0422 .0552 .0571 .0695 .0763 .0857 .0954	370 42 26 6	608 200 88 7	593 16 47 5	836 43 150 15	1250 800 310 330 140 95	0.24 .25 .30 1.8 .86 3.3	1033 760 260 131	33 52 23 33
2050° F, 2 hr, W.Q. + 15% hot- cold-work at 1200° F	10,000 11,500 14,000 18,000 20,000	0.0515 .0747 .0959 .116 .136	6	380 53 8 	^b 56 5.5	10 3	590 143	3.0 3.0	739 321	6 10

aw.Q. - Water-quenched.

Bastimated by creep data extrapolation.

TABLE XIV. - TIME - TOTAL-DEFORMATION STRENGTHS AT 12000 F FOR LOW-CARBON N-155 ALLOY BAR STOCK FROM HEAT 30276

Treatment	Total deformation	Stress (psi) to cause total deformation in -						
(a)	(percent)	10 hr	100 hr	1000 hr	2000 hr			
As-rolled	0.1	21,000	15,500	10,500	9,000			
ST+Ag(1350)	.1	19,000	17,800	14,100	b12,800			
ST+25%HCW	.1	20,000	18,200	16,700	16,000			
As-rolled	.2	29,500	19,800	15,000	13,600			
ST+Ag(1350)	.2	33,500	23,500	19,000	18,000			
ST+25%HCW	.2	37,000	29,800	22,600	20,600			
As-rolled	.5	36,500	30,200	20,000	18,700			
ST+Ag(1350)	.5	37,500	31,300	25,000	23,200			
ST+25%HCW	.5	54,500	45,000	36,000	33,000			
As-rolled	1.0	41,000	33,200	23,600	20,500			
ST+Ag(1350)	1.0	41,000	34,900	28,700	^b 26,900			
ST+25%HCW	1.0	64,500	61,500	54,000	50,000			
As-rolled	Transition		47,500	34,000	¹ 30,000			
ST+Ag(1350)	Transition		42,300	37,000	35,500			
ST+25%HCW	Transition		66,000	54,000	50,000			

Treatments: ST+Ag(1350) - 2200° F, 1 hr, water-quenched; 1350° F, 50 hr. ST+25%HCW - 2050° F, 2 hr, water-quenched; 25% hot-cold-work at 1200° F.

bEstimated.

TABLE XV.- TIME - TOTAL-DEFORMATION STRENGTHS AT 1200°, 1350°, AND 1500° F FOR LOW-CARBON N-155 ALLOY BAR STOCK FROM HEAT A1726

	Total	Str	ess (psi) to cause	total deformation	in -
Treatment (a)	deformation (percent)	10 hr	100 hr	1000 hr	2000 hr
		1200° F			
As-rolled	0.1	19,500	17,200	14,600	b14,000
ST(2100)	.1	1		•	1
ST+Ag(1400)	.1) i			1
811+15% ECW	.1	1			
As-rolled	.2	35,000	26,800	18,700	b16,500
BT(2100)	.2	30,500	27,000	h	
ST+Ag(1400)	.2	32,400	26,000	^b 19,500 ^b 25,000	b22,500
ST+15%HCW	••	38,300	32,300	-27,000	-22,000
As-rolled	.5	12,000	34,800	27,300	b25,000
BP(2100)	•5 •5 •5	36,100	34,700	23,500	b18,500
BT+Ag(1400) BT+15%HCW	•2	38,000 52,000	32,000	25,500	23,600
DITTEN	• • • • • • • • • • • • • • • • • • • •	/2,000			,
As-rolled	1.0	¥7,000	39,500	31,700	h29,500
BT(5100)	1.0	38,500	36,100	30,700	26,500
ST+Ag(1400) ST+15%HCW	1.0 1.0	40,500	34,700 53,000	28,800	b27,000
· · · · · · · · · · · · · · · · · · ·					1
As-rolled	Transition		^{Ֆլ} 47,000	har eee	1
ST(2100) ST+Ag(1400)	Transition Transition		45,800	^b 37,000 31,500	b25,000
SI+15KHCW	Transition		⁵ 5,000	31,700	-27,000
		1350° I		<u> </u>	
		1		N 144	r
As-rolled ST(2100)	0.1 .1	14,200	10,800 b9,500	7,400	1
BT+Ag(1400)	.1	15,400	11,000	¹6,500	1
SII+15% HCW	.1		p11,300	,,	
	•		16 600	33.000	0 100
As-rolled ST(2100)	.2 .2	22,000	16,600 14,300	11,000 18,200	9,400
ST+Ag(1400)	.2	21,500	16,000	10,500	8,900
BT+15%HCW	.2	27,500	21,700	15,800	14,000
As-rolled	R	28,700	22 200	16 000	b14,000
8T(2100)	.5	24,300	22,300 18,800	16,000 13,300	11,800
BT+Ag(1400)	.5 .5 .5	1 24.200 . 1	20,100	16,000	14,800
ST+15%HCW	.5	37,500	30,600	24,000	22,000
As-rolled	1.0	32,500	26,000	19,600	b17,700
ST(2100)	LO.	27,000	21,300	16,900	15,500
ST+Ag(1400)	1.0	27,300	22,200	₱17,200	
8T+15≰HCW	1.0	41,000	34,000	27,000	b24,800
As-rolled	Transition		32,200	³28,000	
BT(2100)	Transition		26,700	22,500	P20,500
BT+Ag(1400)	Transition		29,500 38,500	⁰ 21,300 26,500	
BT+15KHCW	Transition			26,500	23,000
		1500°	P		
As-rolled	0.1	10,100	6,800	b3,500	
8T(2100)	• 1	10,500	7,000 8,500	5,800	De 000
ST+Ag(1400) ST+15%HCW	.1	11,300 b9,200	8,500	3,000	000,5م
		,,		1	
As-rolled	.2	12 000	10,000	6,600	05,500 05,000
ST(2100) ST+Ag(1400)	.2 .2	13,200 15,100	10,800 11,600	6,500 8,000	97,000
BIT+15KHCW	.2	13,700	11,200	9,000	b8,000
1			,		
As-rolled	٠ <u>٠</u>	11.200	p12,800	^ъ 9,700 ^ъ 10,200	
BT(2100) ST+Ag(1400)	•5 ••5	14,200	15,000	b11,000	^b 9,500
ST+15XECW	.5	17,000	b13,200]	1
4e-molled	1.0	13.000			
As-rolled ST(2100)	1.0 1.0	13,000 15,100	b13,700	b11,700	
BT+Ag(1400)	1.0	18,700	16,300	11,500	b10,000
ST+15%HCW	1.0	18,000	b14,200	1	'
As-rolled	Transition		14,500	9,500	₽8,000
BT(2100)	Transition			}` ¤8,000	,,,,,,
ST+Ag(1400)	Transition		17,200	9,600	7,500
ST+15/CHCW	Transition		20,500		

STreatments: ST(2100) - 2100° F, 1 hr, water-quenched. ST+Ag(1400) - 2200° F, 1 hr, water-quenched; 1400° F, 24 hr. ST+15%HGW - 2050° F, 2 hr, water-quenched; 15% hot-cold-work at 1200° F.

TABLE XVI.- CREEP DATA AT 1200° F FOR LOW-CARBON N-155 ALLOY BAR STOCK FROM HEAT 30276

Mara day and	8tress	Duration	Initial deformation (percent)	Creep rate (percent/hr) at -					
Treatment	(psi)	(hr)		500 hr	1000 hr	1500 hr	M inimm ⁸		
As-rolled	10,500	1290	0.044	0,000050	0.000021				
	11,000	1475	.044	.000045	.000029	0.000029	į		
	12,500	1030	.048	.000062	.000053		}		
	13,500	1270	.061	.000083	.000058	ŀ	1		
	15,000	1406	.066	.000096	.000067	.000067	Ì		
	18,000	3430	.076	.000200	.000155	.000135	·		
	20,000	1004	.100	.000318	.000268		1		
	22,500	1,006	.128	,000600	.000480	j	l		
	25,000	1000	.128	.00090	.000615	Ţ	ļ		
	30,000	1400	.148	.00166	.00105	.00105			
	40,000	b360	.331				0.015		
	45,000	ъ150	1.26				.038		
2200° F, 1 hr, W.Q.°	15,000	1415	0.066	0.000027	0.000019	0.000019			
+ 50 hr at 1350° F	17,500	1265	.074	.000034	.000026				
1 2	20,000	1010	.096	.000120	.000100				
	25,000	1004	.123	.000275	.000220		ł		
	30,000	1004	.14 5	.00110	.000880		ŀ		
	35,000	^b 2700	.221	.00320	.00350	.00350	0.0032		
	40,000	p ₂₉₀	.386				.0100		
	45,000						.0205		
2050° F, 1 hr, W.Q.	15,940	1818	0.062	0.0000105	0.0000085	0.0000085			
+ 25% hot-cold-work	20,000	1372	.092	.0000380	.0000380		1		
at 1200° F	30,000	1868	.138	.000105	.000080	.000058			
•	40,000	,1218	.169	.000380	.000110		1		
	55,000	b900	.232				0.00038		
	60,000	b425	-253				.00165		
	65,000	b ₁₂₅	.274				.0035		

^aMinimum rates of tests entering third-stage creep. ^bFine of transition to third-stage creep.



CW.Q. - Water-quenched.

Table XVII.- Creep data at 1200°, 1350°, and 1500° F for low-carbon H-155 ALLOY BAR STOCK FROM HEAT A1726

Treatment	Stress	Duration			Creep rate (pe	rcent/hr) at -	~	
Treatment	(psi)	(hr)	(percent)	500 hr	1000 hr	1500 hr	Kinimm ^a	
·			1200° F					
As-rolled	15,000 20,000 25,000 30,000 32,500 42,000	1630 1534 690 1586 863 9375	0.0550 .0905 .1162 .1273 .1432	0.000023 .000099 .000175 .00039 .00093	0.000023	.00025	0.0081	
,	45,000 47,000	0190 115	.196				.0137	
2100° F, 1 hr, W.Q.°	27,500 30,000 32,500 35,000 37,000 40,000	1238 1268 1238 982 5920 5720	0.120 .160 .225 .320 .430	0.000700 .000850 .000930 .001 <i>6</i> 8	0.000390 .000470 .000560 .00157		0.00225 .0048b	
2200° F, 1 hr, W.Q. + aged 24 hr at 1400° F	25,000 30,000 34,000 44,000 46,000	1675 1150 1955 1955 1914	0.1125 .1366 .1759 .636	0.00025 .00114 .00500	0.00025	0.00025	.00214 .0120 .0360	
2050° F, 2 hr, W.Q. + 15% hot-cold-work at 1200° F	30,000 37,000 42,000 48,000 49,000 52,000	1458 1310 257	0.1360 .1690 .1900 .217 .222	0.000033	0.000033	0,000033	0.000332 .00075 .00070	
	55,000		.249				00200	
			13500 1				Γ	
As-rolled	10,000 15,000 20,000 30,000 32,500	1657 1651 1567 1335 169	0.0467 .0769 .0957 .1810 .240	0.00096 .000200 .000192	0.000054 .000111 .000357	0.00030 .00056 .000377	0.0082	
2100° F, 1 hr, W.Q.	12,500 15,000 17,500 20,000 25,000 27,000 29,000 30,000	1464 1274 1464 1263 8405 8220 825 865	0.0530 .0630 .0740 .0650 .095 .112 .135	0.000185 .000325 .000378 .001075	0.000072 .000137 .000195 .000750	0.000037	0.0138 .0360 .0540 .0730	
2200° F, 1 hr, W.Q. + aged 24 hr at 1400° F	10,000 15,000 17,500 20,000 26,000 30,000	1657 1483 1315 864 ^b 270 ^b 90	0.0483 .0731 .0781 .1024	0.000095 .000197 .000303 .000760	0.000020 .000064 .000135 .000690	0.000020 .000048 .000065	0.0290 .0510	
20500 F, 2 im, W.Q. + 15% hot-cold-work at 12000 F	15,000 20,000 26,500 25,000 33,000 36,000 38,000	1561 1127 1555 12410 1312 195 195	0,0690 .0910 .1030 .1160 .149 .162	0.000089 .000133 .000123 .000343	0.000048 .000133 .000123 .000343	0.0000\(\frac{1}{2}\) .000123 .000370	0.000343 .0026 .0037 .0090	
			1590° F			·		
As-rolled	7,500 9,000 10,500 12,000 13,000 13,500	1215 91225 450 9400 9110 9260 972	0.0340 .0442 .0564 .0580 .0630 .0680	.000206 .000200 .000400	.000200	0.000360	0.000200 .0032 .0120 .0140	
2100° F, 1 hr, W.Q.	8,500 10,000 12,000 15,000 16,500	1152 1278 0410 0234 0136	0.0409 .0433 .0546 .181 .240	0.000086 .000063 .00099	0.000013		0.000446 .0114 .0560	
2200° F, 1 hr, W.Q. + aged 24 hr at 1400° F	7,000 9,000 10,000 12,000 14,600 16,000 18,000	1483 51250 5800 5310 5330 5140	0.0328 .0422 .0552 .0571 .0695 .0763 .0857	0.000028 .000084 .000069 .000785	0.000019 .000052 .000105 .00494	0.000010	0.00052 .00069 .000350 .00210 .00390	
2050° F, 2 hr, W.Q. + 15% hot-cold-work at 1200° F	10,000 11,500 12,500 18,000 20,000	1119 1000 1274 5424 5143	0.0515 .0747 .0595 .116 .136	0.000101 .000132 .000099	0.000081 .000089 .000074		0.00290 .00625	

*Minimum rates of tests entering third-stage creep. brime for transition to third-stage creep. CW.Q. - Water-quenched.



TABLE XVIII. - CREEP STRENGTHS OF TWO HEATS OF LOW-CARBON N-155 ALLOY BAR STOCK

			Creep strengths (psi) from tests at -					
Treatment	Heat	Temperature (°F)	0,00001	percent/hr	0.0001 percent/hr			
(a)			1000 hr	1500 hr	1000 hr	1500 hr		
As-rolled	30276 A1726	1200 1200	8,200 11,800	11,800	15,500 23,000	23,000		
ST(2100)	A1726	1200						
ST+Ag(1350)	30276	1200	13,700 .		20,800			
ST+Ag(1400)	A1726	1200				22,000		
8T+15≸HCW	A1726	1200	23,500	23,500	37,000	ъ37,000		
ei+25%ecw	30276	. 1200	15,500		34,000			
As-rolled	A1726	1350	ъ4,000	^b 5,000	14,000	16,000		
8T(21.00)	- A1726	1350		ъ9, 5 00	14,000	15,000		
ST+Ag(1400)	A1726	1350	8,000	8,000	16,000	17,500		
9T+15%HCW	A1726	1350	¹⁰ 8,400	b8,400	19,500	19,500		
As-rolled	. A1726	1500			8,200			
ST (2100)	A1726	1500	8,000		11,000			
9T+Ag(1400)	A1,726	1,500	5,800	7,000	10,000			
9T+15%HCW	A1726	1500	=====		11,000			

**BTreatments: ST(2100) - 2100° F, 1 hr, water-quenched.

ST+Ag(1350) - 2200° F, 1 hr, water-quenched; 1350° F, 50 hr.

ST+Ag(1400) - 2200° F, 1 hr, water-quenched; 1500° F, 24 hr.

ST+155kW - 2050° F, 2 hr, water-quenched; 15% hot-cold-work at 1200° F.

ST+256kW - 2050° F, 2 hr, water-quenched; 25% hot-cold-work at 1200° F.

bEstimated.

TABLE XIX.- COMPARATIVE CREEP AND RUPTURE STRENGTHS OF

LOW-CARBON N-155 ALLOY BAR STOCK

Treatment	Heat	Temperature (°F)	Creep strength, 0.0001 percent/hr (psi)	Rupture strength, 10,000 hr (psi)
As-rolled	30276	1200	15,500	28,000
As-rolled	A1726	1200	23,000	38,000
2100° F, 1 hr, W.Q. ⁸ 2200° F, 1 hr, W.Q. + aged 1350° F for 50 hr 2200° F, 1 hr, W.Q. + aged 1400° F for 24 hr 2050° F, 2 hr, W.Q. + 15% hot-cold-work 2050° F, 2 hr, W.Q. + 25% hot-cold-work	A1726	1200	^b 15,000	34,000
	30276	1200	20,800	32,000
	A1726	1200	22,000	38,000
	A1726	1200	37,000	42,000
	30276	1200	34,000	45,000
As-rolled	A1726	1350	14,000	24,000
2100° F, 1 hr, W.Q.	A1726	1350	14,000	20,000
2200° F, 1 hr, W.Q. + aged 24 hr at 1400° F	A1726	1350	16,000	20,000
2050° F, 2 hr, W.Q. + 15% hot-cold-work	A1726	1350	19,500	23,000
As-rolled	A1726	1500	8,200	8,600
2100° F, 1 hr, W.Q.	A1726	1500	11,000	11,500
2200° F, 1 hr, W.Q. + aged 24 hr at 1400° F	A1726	1500	10,000	10,000
2050° F, 2 hr, W.Q. + 15% hot-cold-work	A1726	1500	11,000	12,000

W.Q. - Water-quenched.

bEstimated.

TABLE IX. - EFFECT OF CREEF TERFING AT 1200°, 1350°, AND 1500° F ON PHYSICAL PROPERTIES AT BOOM TEMPERATURE OF LOW-CARBON M-155 ALLOY BAR STOCK FROM HEAT A1726

	Test	ing conditio	ne	Tensile	Offse	t yield Str (psi)	rength	Propor-	Klonga-	Reduc-	Izod	We also
Treatment	Temper- ature (°F)	Stress (psi)	Time (hr)	strength (psi)	0.02 percent	0.1 percent	0.2 percent	tional limit (psi)	tion in 2 in. (percent)	tion of area (percent)	impact strength (ft-lb)	Vickers hardness
As-rolled		Original		127,200	63,000	73,200	77,400	46,500	40	51	101, 97	249
,	1200 1200	15,000 32,500	1680 863	129,000	65,300 	75,800 	79,800	53,000	34 ~-~-	38.5	26, 28	262
	1350 1350	10,000 · 20,000	1657 1567	131,000	62,200	75,700	79,200	43,500 	18 	16	. B, 8	282
	1500 1500	7,500 10,500	1215 450	128,000	53,000	64,800	70,100	39,500	27 —	27.5	12, 12	261
2100° 7, 1 hr,		Original		117,000	45,700	52,800	56,300	36,000	50	67.6	116, 115	201
W.Q. ¹	1200 1200	27,500 30,000	1238 1268	127,000	66,000	72,100	74,700	55,500	33.5	36 	33, 35	258
	1350 1350	12,500 20,000	1464 1268	137,500	61,500	70,000 	74,700	50,000	14.5 	12.5	11, 11	280
	1500 1500	8,500 12,000	1152 1100	115,000	43,700	53,700	58,800	33,000	11.5	10.8	15, 18	233
2200° F, 1 hr,		Original		118,900	47,800	55,800	59,400	37,000	38	46.8	33, 33	231
W.Q. + aged 24 hr at : 1400° F	1200 1200	25,000 34,000	1675 495	125,500	60,400	68,400.	71,800	52,000	21	18.5	8, 8	263
1	1350 1350	10,000 20,000	1484 864	130,000	56,700	69,500	74,700	39,500	13.5	11.7	4, 4	282
•	1500 1500	7,000 12,000	1483 1079	116,000	¥9,300	59,100	64,200	38,000	13 	10	6, 7	251
2050° F, 2 hr,		Original		144,000	107,500	117,200	120,300	97,500	24	57.0	76, 71	313
W.Q. + 15% hot-cold-work at 1200° F	1,200 1200	30,000 37,000	1458 1310	141,000	97,700	108,700	113,000	85,000	20.5	. 33.8		313
	1350 1350	15,000 25,000	1561 1639	136,500	75,000	88,200	93,900	58,000 	8	13.5	3, 5	 292
	1500 1500	10,000 11,500	1119 1000	127,500	64,000	77,200	83,700	46,000	14.5	15.6 	8, 8	<u>261</u>

lw.Q. - Water-quenched.

NACA

TABLE XXI.- LOW-ANGLE X-RAY DIFFRACTION PARTERNS FROM RUPTURE SPECIMENS OF SOLUTION-THEATED AND HOT-COLD-WORKED 25-PERCENT LOW-CARBON N-155 ALLOY FROM HEAT 30276

Cr Ko rediction

Di	ffraction patterns of low-carbo	from rupture sg m H-155 alloy	oecimens	Standard diffraction patterns						
	1500° F - 13,000 psi - 340 hr		30,000 psi - Sigma phas A6 hr (reference		Signa phase (reference 4) CbC (refere		eference 5)	Indicated phase in low-carbon N-155		
d (A)	Intensity (1)	d (A)	Intensity (1)	д (A)	Intensity (1)	đ (A)	Intensity (1)			
2.52	н	2.54	Ж	F		2.54	В	СРС		
			er re	2,50 2,360 2,323 2,268	AM AM AM AM					
2.182	¥	2.21	VW			2.20	В	СрС		
2.12 	W(M) W W W	1.97	M W W	2.115 2.067 2.034 2.020 1.968 1.926 1.882 1.836	W W W B · B W			81gma 81gma Sigma 81gma Sigma		
1.556 1.32	8 8	1.56 1.327	H	1.326	N	1.55 1.33	s N	CbC CbC and Signa		
1.248 1.23 1.221 1.21 1.20	¥ ¥ ¥ ¥			1.295 1.256 1.240 1.235 1.229 1.224 1.218 1.205 1.198 1.186	VW W W W W M M M M W W W W W W W W W W W			Sigma Sigma Sigma Sigma Sigma		
1.176	x(s)			1.175 1.166	W(M) M(S)	'	,	Bigus		

 $^{^{1}}$ M, medium; W, weak; S, strong; \forall , very.



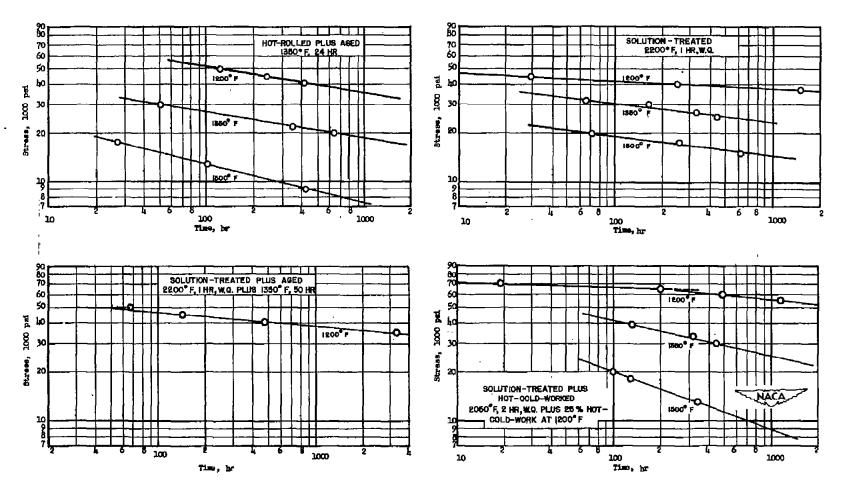
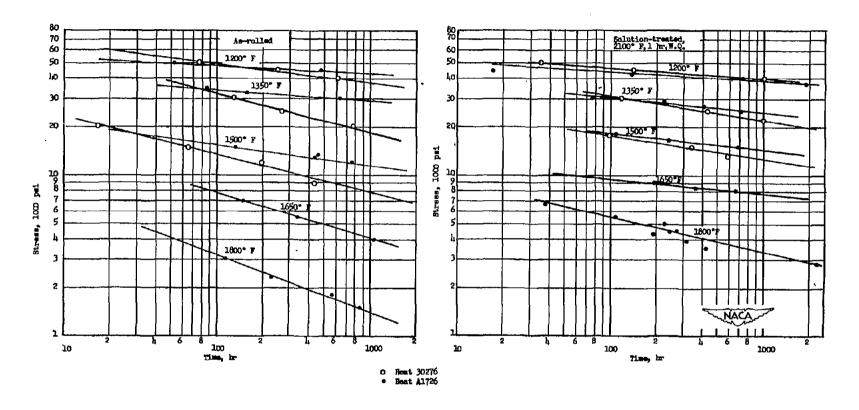
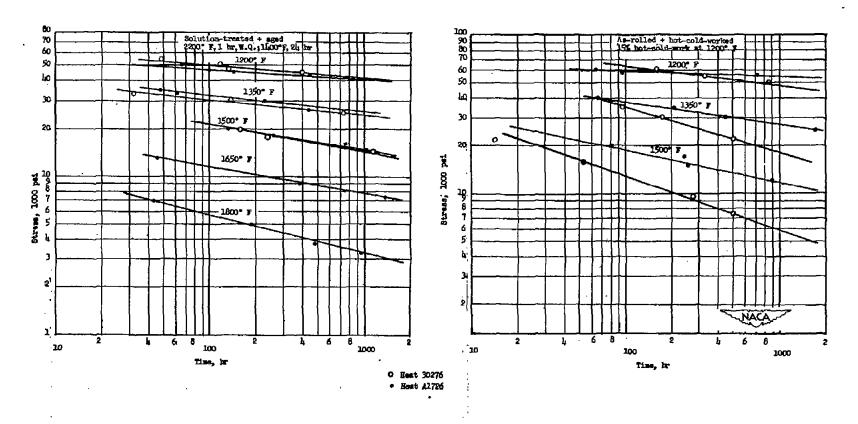


Figure 1.- Curves of stress against rupture time at 1200°, 1350°, and 1500° F for low-carbon N-155 alloy bar stock from heat 30276.



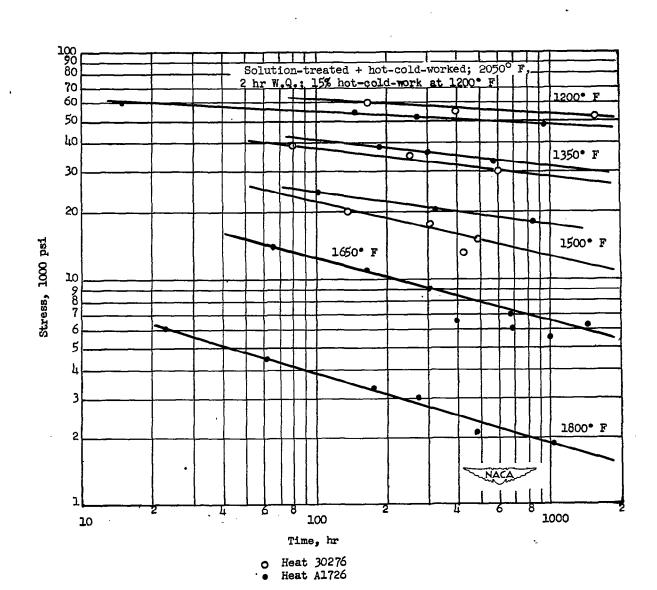
(a) As-rolled and solution-treated conditions.

Figure 2.- Curves of stress against rupture time at 1200° to 1800° F for low-carbon N-155 alloy bar stock from two heats.



(b) Solution-treated plus aged and as-rolled plus hot-cold-worked conditions.

Figure 2.- Continued.



(c) Solution-treated plus hot-cold-worked condition.

Figure 2.- Concluded.

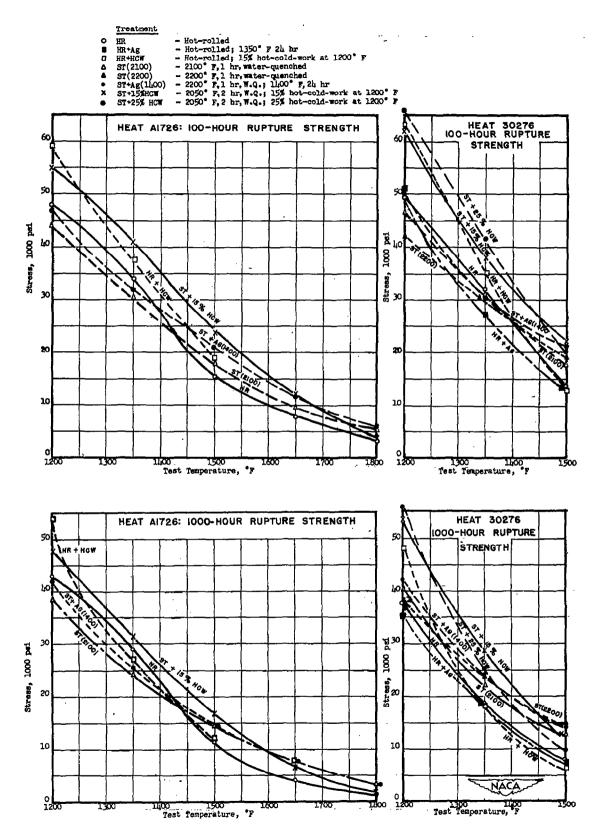


Figure 3.- Influence of testing temperature on rupture strength of low-carbon N-155 alloy bar stock from two heats.

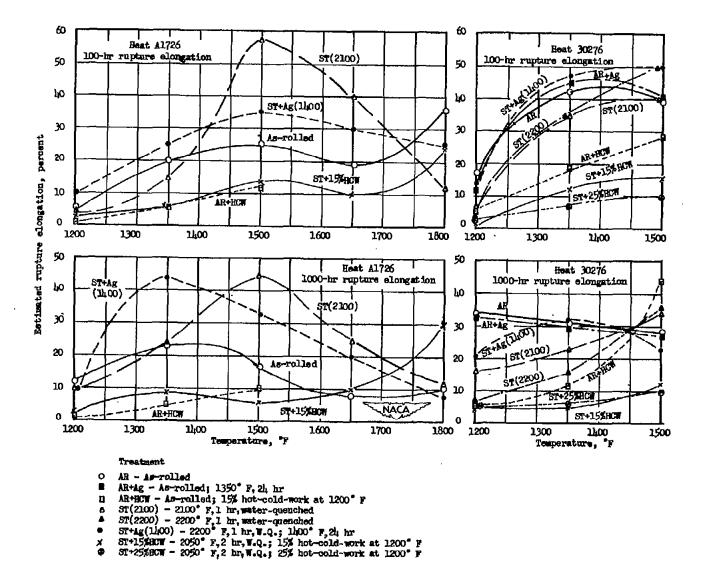
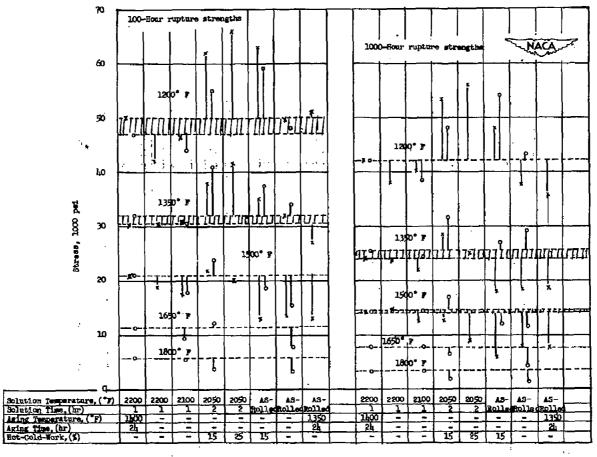


Figure 4.- Influence of testing temperature on rupture elongation of low-carbon N-155 alloy bar stock from two heats.



O Heat al726 X Heat 30276

Figure 5.- Rupture strengths of variously treated low-carbon N-155 bar stock compared with rupture strengths of solution-treated and aged condition. Horizontal dashed lines indicate range of properties for the two heats in the solution-treated and aged condition (2200° F, 1 hr, water-quenched; 1400° F, 24 hr).

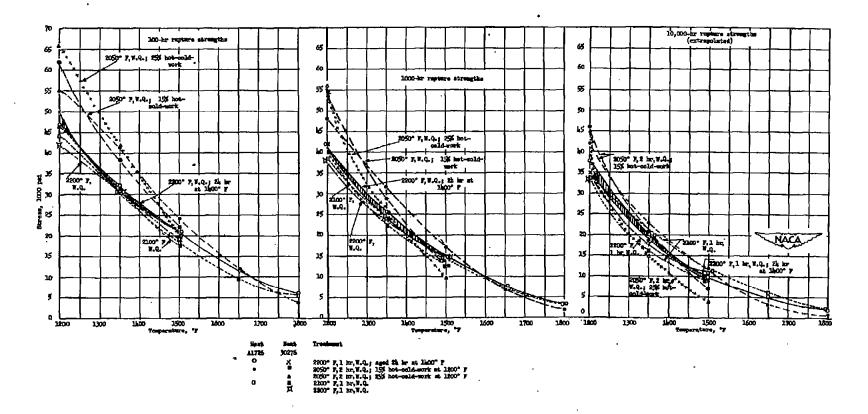


Figure 6.- Influence of temperature on relative rupture strengths of solution-treated and aged, solution-treated and hot-cold-worked, and solution-treated low-carbon N-155 alloy bar stock. Shaded area indicates property ranges for two heats in solution-treated and aged condition (2200° F, 1 hr, water-quenched; 1400° F, 24 hr).

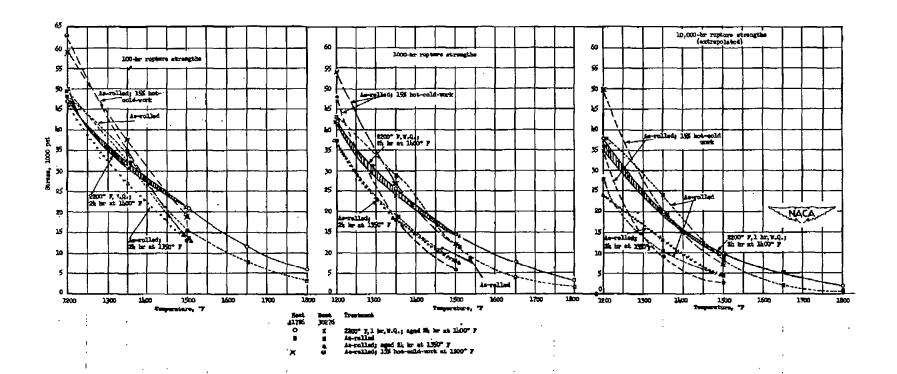
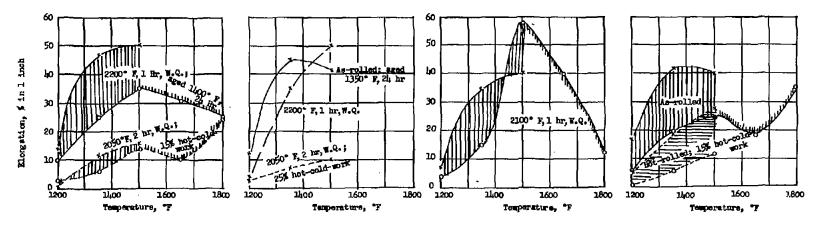
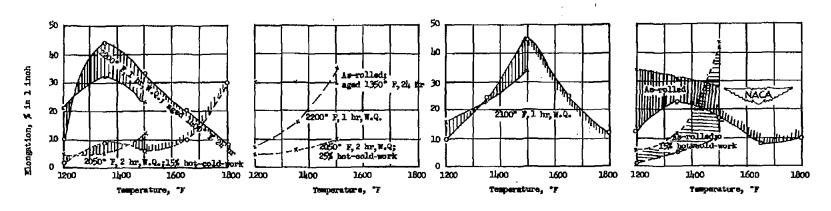


Figure 7.- Influence of temperature on relative rupture strengths of solution-treated and aged, as-rolled, as-rolled and aged, and as-rolled and hot-cold-worked low-carbon N-155 alloy bar stock. Shaded area indicates property ranges for two heats in solution-treated and aged condition (2200° F, 1 hr, water-quenched; 1400° F, 24 hr).



(a) Estimated elongation for rupture in 100 hours.



o Heat 11726

(b) Estimated elongation for rupture in 1000 hours.

Figure 8.- Influence of testing temperature and prior treatment on rupture elongations of low-carbon N-155 alloy bar stock from two heats.

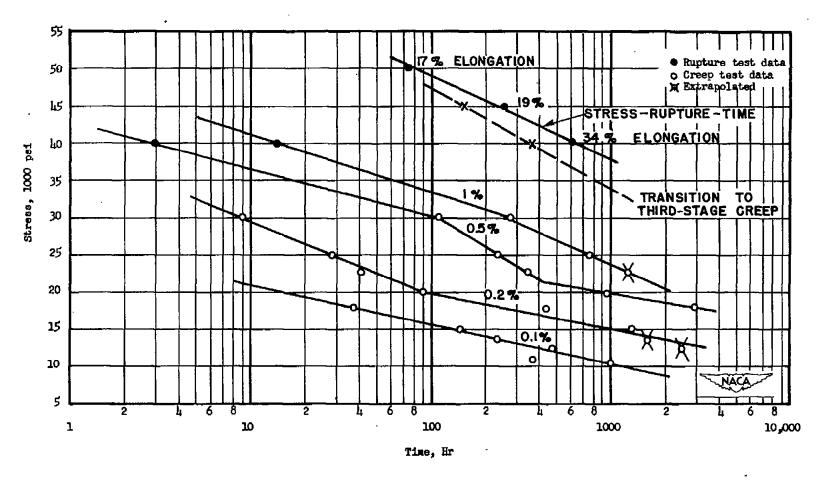


Figure 9.- Curves of stress against time for total deformation at 1200° F for hot-rolled low-carbon N-155 alloy bar stock from heat 30276.

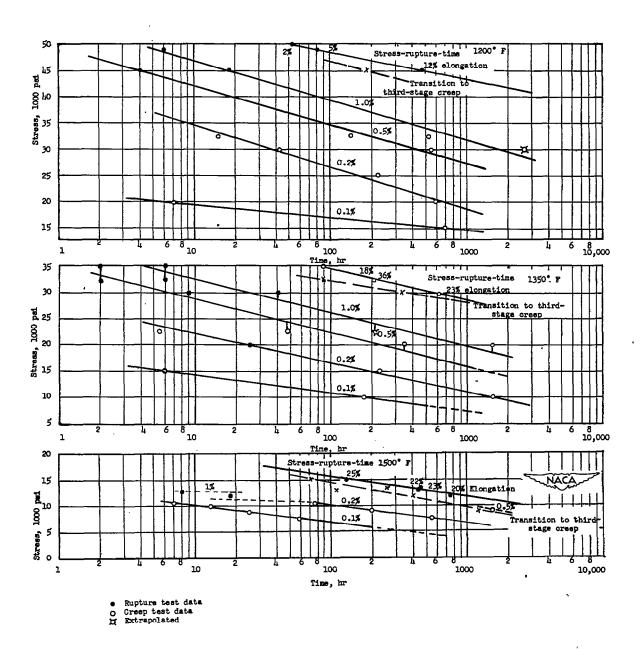


Figure 10.- Curves of stress against time for total deformation at 1200°, 1350°, and 1500° F for hot-rolled low-carbon N-155 alloy bar stock from heat A1726.

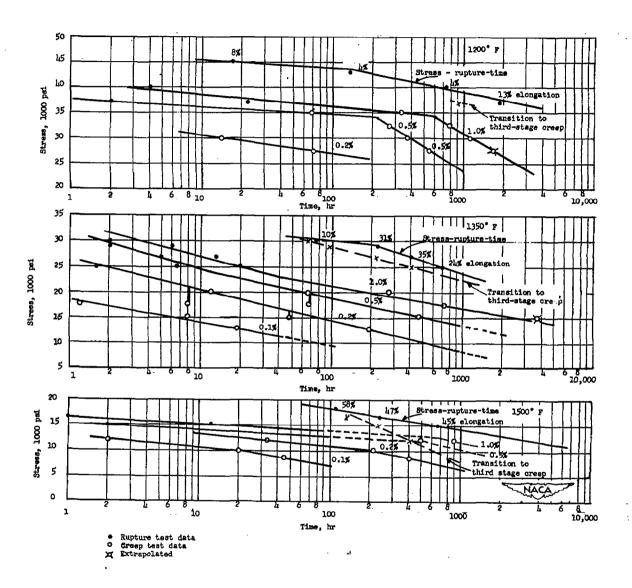


Figure 11.- Curves of stress against time for total deformation at 1200°, 1350°, and 1500° F for solution-treated low-carbon N-155 alloy bar stock from heat A1726. (Treatment: 2100° F, 1 hr, water-quenched.)

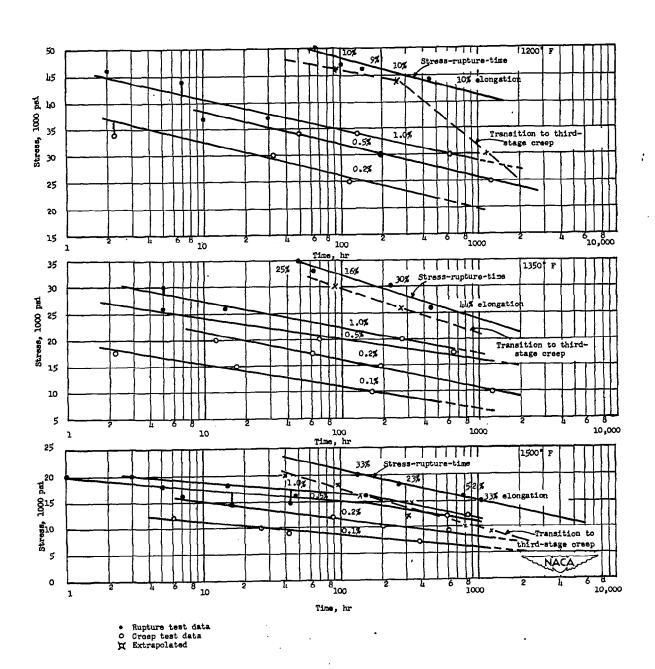


Figure 12.- Curves of stress against time for total deformation at 1200°, 1350°, and 1500° F for solution-treated and aged low-carbon N-155 alloy bar stock from heat A1726. (Treatment: 2200° F, 1 hr, water-quenched; 1400° F, 24 hr.)

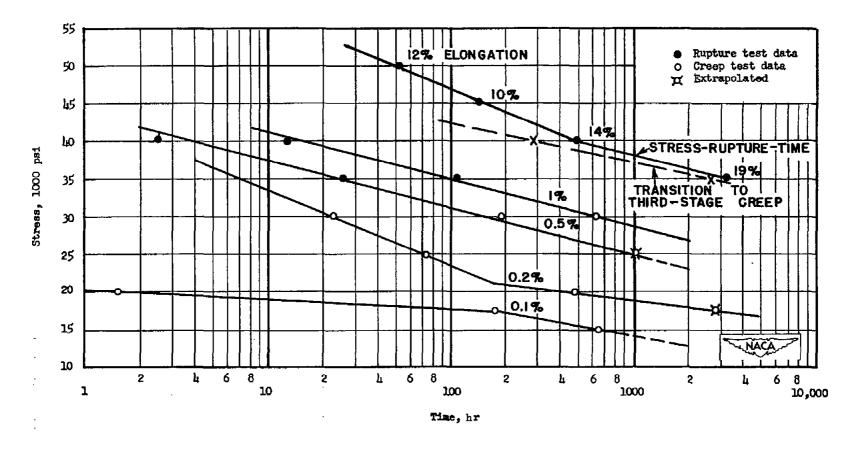


Figure 13.- Curves of stress against time for total deformation at 1200° F for solution-treated and aged low-carbon N-155 alloy bar stock from heat 30276. (Treatment: 2200° F, 1 hr, water-quenched; 1350° F, 50 hr.)

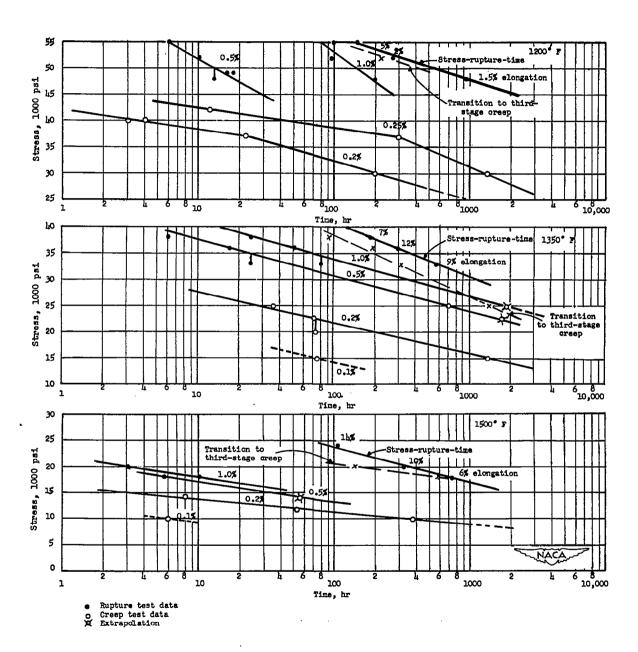


Figure 14.- Curves of stress against time for total deformation at 1200°, 1350°, and 1500° F for solution-treated and hot-cold-worked low-carbon N-155 alloy bar stock from heat A1726. (Treatment: 2050° F, 2 hr, water-quenched; 15-percent hot-cold-work at 1200° F.)

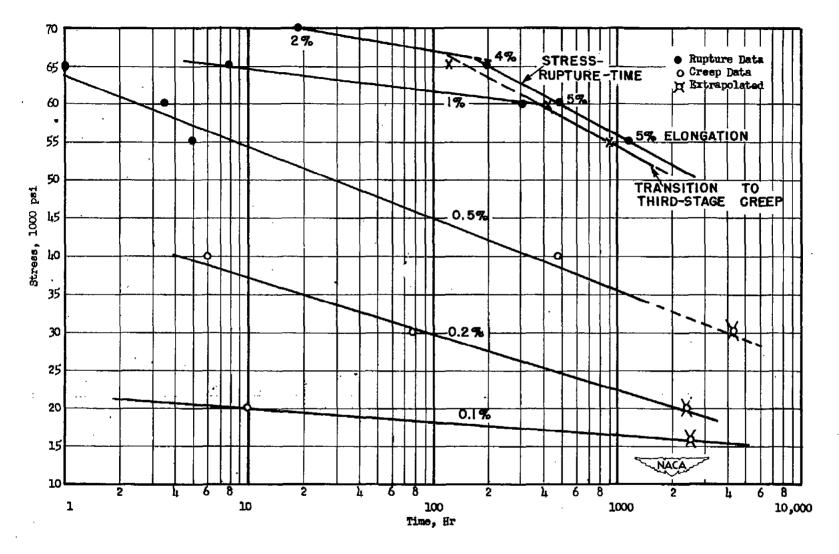


Figure 15.- Curves of stress against time for total deformation at 1200° F for solution-treated and hot-cold-worked low-carbon N-155 alloy bar stock from heat 30276. (Treatment: 2050° F, 2 hr, water-quenched; 25-percent hot-cold-work at 1200° F.)

NACA RM 51B05

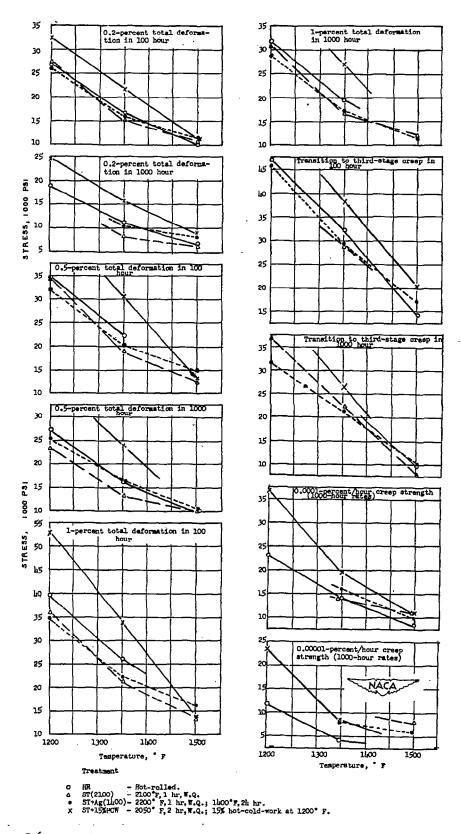


Figure 16.- Influence of testing temperature on total-deformation strengths and creep strengths of low-carbon N-155 alloy bar stock from heat Al726.

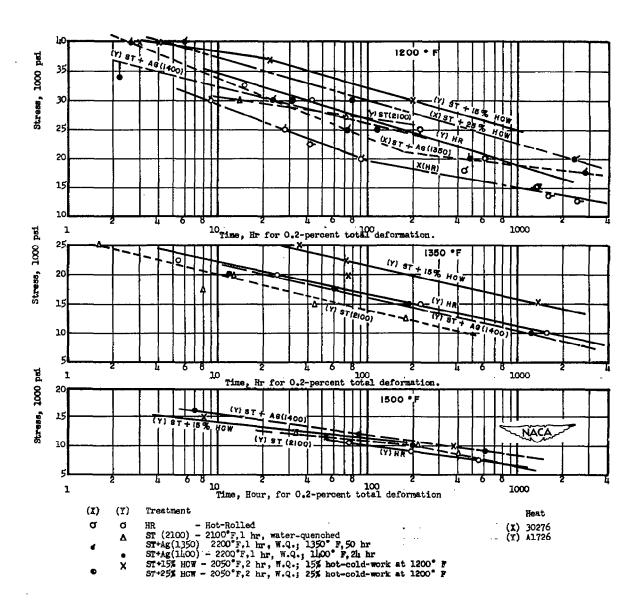


Figure 17.- Comparative 0.2-percent total-deformation characteristics at 1200°, 1350°, and 1500° F for low-carbon N-155 alloy bar stock from two heats.

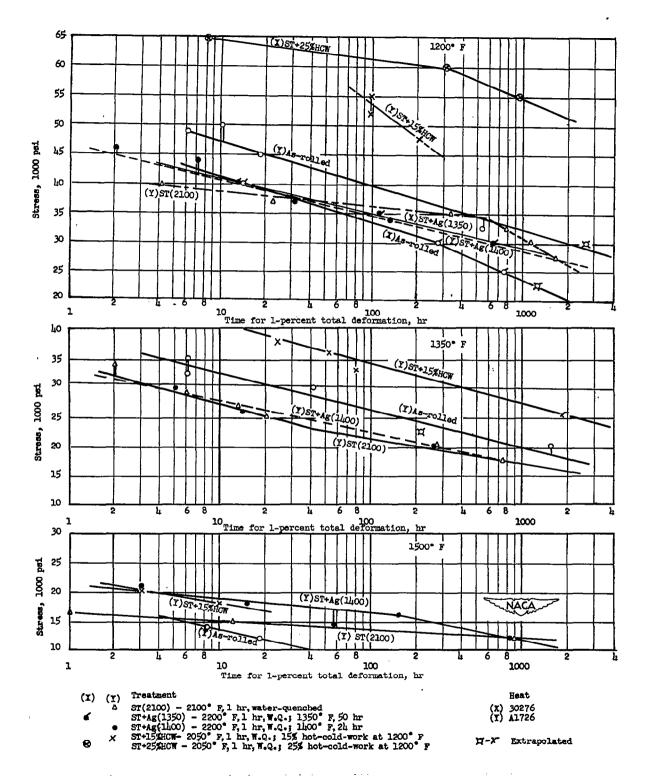
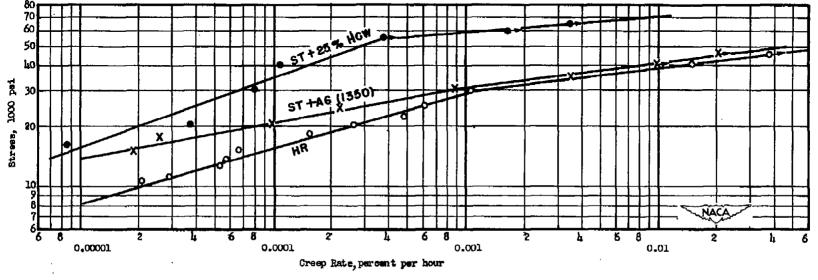


Figure 18.- Comparative 1-percent total-deformation characteristics at 1200°, 1350°, and 1500° F for low-carbon N-155 alloy bar stock from two heats.



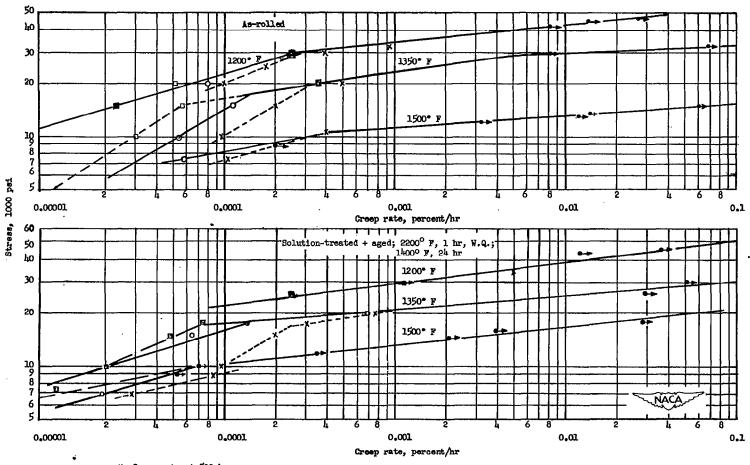


- Hot-Rolled

X ST+Ag (1350) - 2200°F, 1 Hr, W.Q.; plns 1350°F, 50 Hr. • ST+25% HCW - 2050°F, 2 Hr, W.Q.; plns 25% hot-cold-work at 1200°F

- Third-stage creep occurred.

Figure 19.- Curves of stress against creep rate at 1200° F for low-carbon N-155 alloy bar stock from heat 30276. Rates shown are those at 1000 hours or minimum rates if third-stage creep occurred.



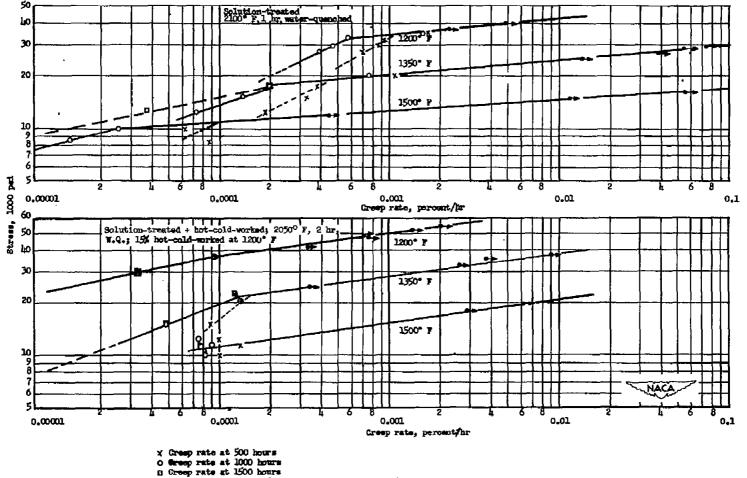
X Greep rate at 500 hours O Greep rate at 1000 hours

(a) As-rolled and solution-treated plus aged conditions.

Figure 20.- Curves of stress against creep rate at 1200°, 1350°, and 1500° F for low-carbon N-155 alloy bar stock from heat A1726.

O Creep rate at 1500 hours

Minimum creep rates if third-stage creep occurred



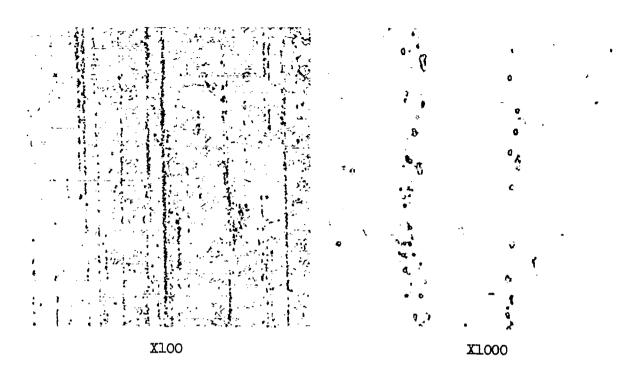
- --- Ministen creep rates if third-stage creep occurred

(b) Solution-treated and solution-treated plus hot-cold-worked conditions.

Figure 20.- Concluded.

NACA RM 51B05

Э



(a) Heat Al726 - as-rolled.

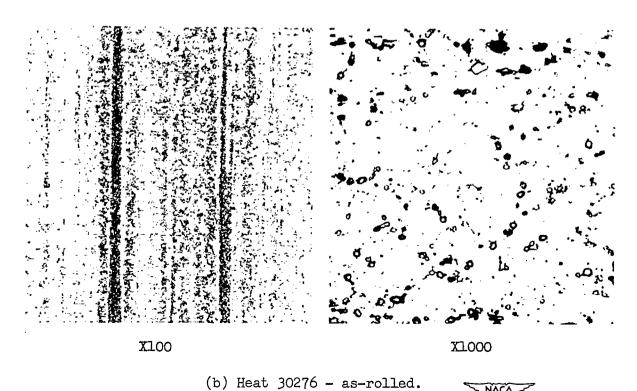
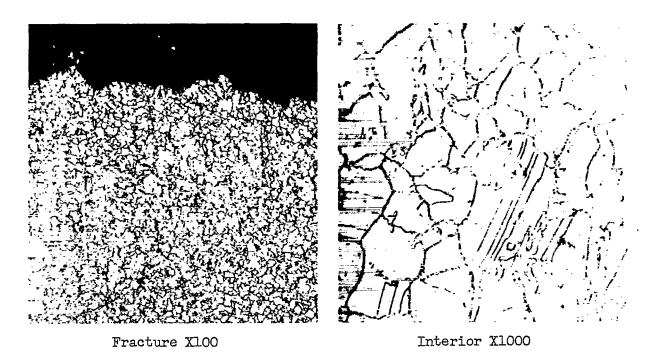
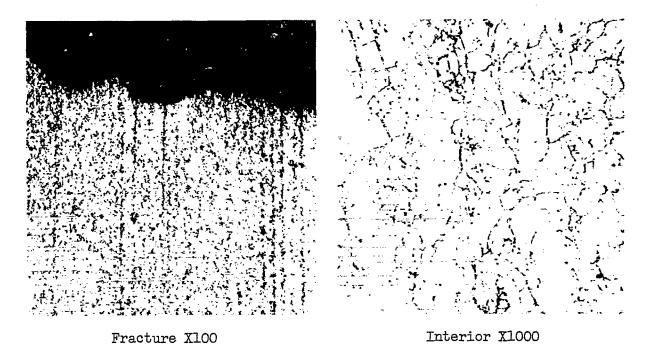


Figure 21.- Original microstructures and microstructures after rupture testing of as-rolled low-carbon N-155 alloy bar stock from heats A1726 and 30276. Electrolytically etched in 10 percent chromic acid.

*
-



(c) Heat Al726 - ruptured in 472 hours under 45,000 psi at 1200° F.



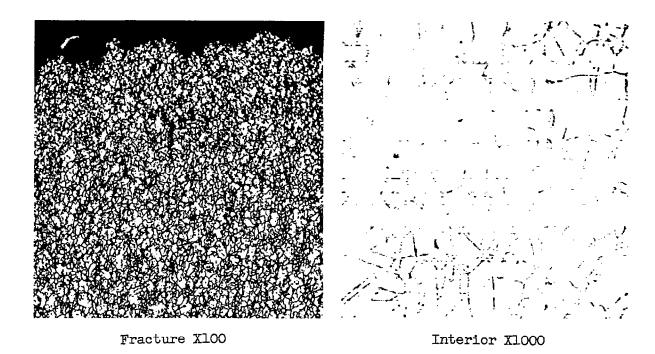
(d) Heat 30276 - ruptured in 610 hours under 40,000 psi at 1200° F.

Figure 21.- Continued.

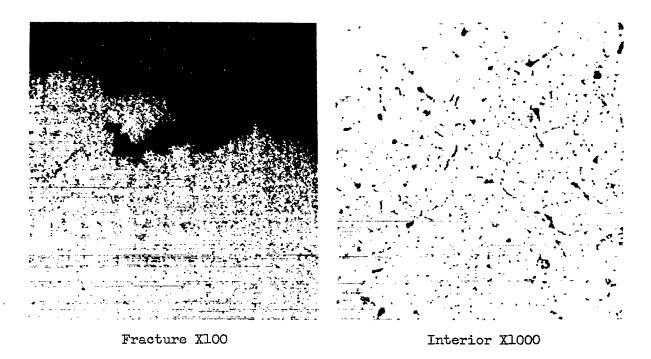
	٠
	•
	•
•	
	òs

65

11



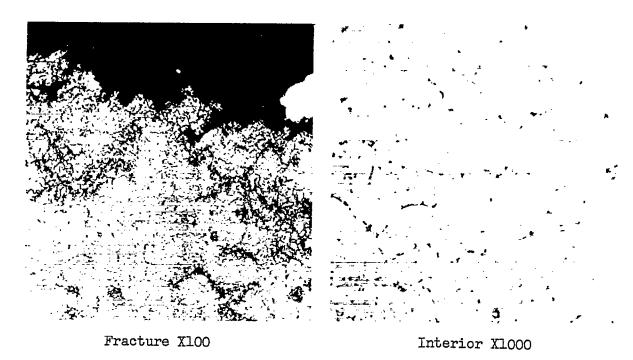
(e) Heat Al726 - ruptured in 623 hours under 30,000 psi at 1350° F.



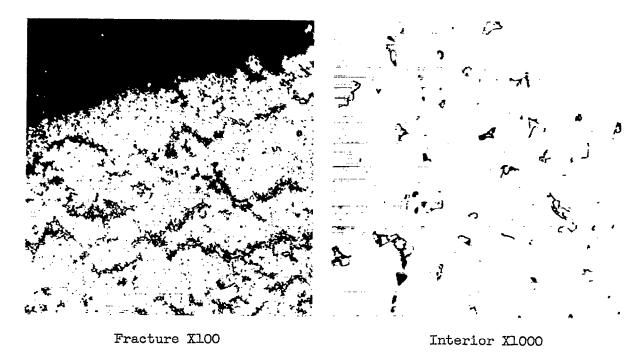
(f) Heat 30276 - ruptured in 130 hours under 30,000 psi at 1350° F. Figure 21.- Continued.

			•
·			
			u
			•
		 ·	

NACA RM 51B05



(g) Heat Al726 - ruptured in 747 hours under 12,000 psi at 1500° F.



(h) Heat 30276 - ruptured in 430 hours under 9000 psi at 1500° F. Figure 21.- Continued.

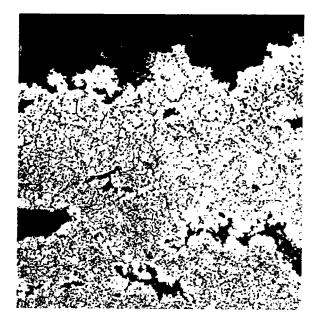
.2



Fracture X100

Interior X1000

(i) Heat Al726 - ruptured in lo51 hours under 4000 psi at 1650° F.





Fracture X100

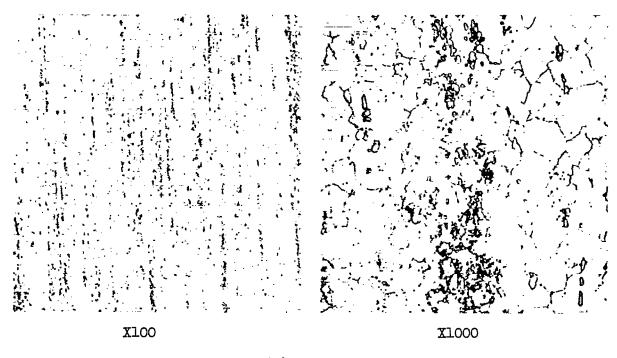
Interior X1000

(j) Heat Al726 - ruptured in 847 hours under 1500 psi at 1800° F.

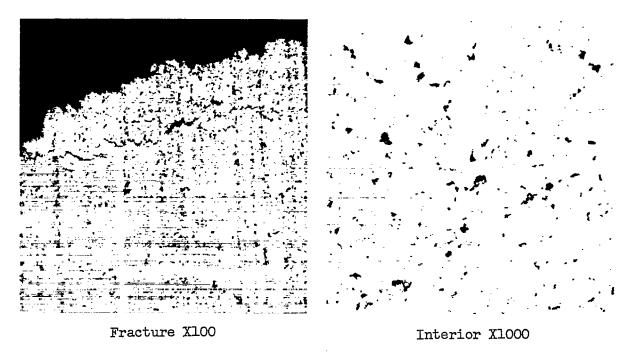
Figure 21.- Concluded.

			ě
			•
			•
•			
			~
			•

3



(a) Initial.

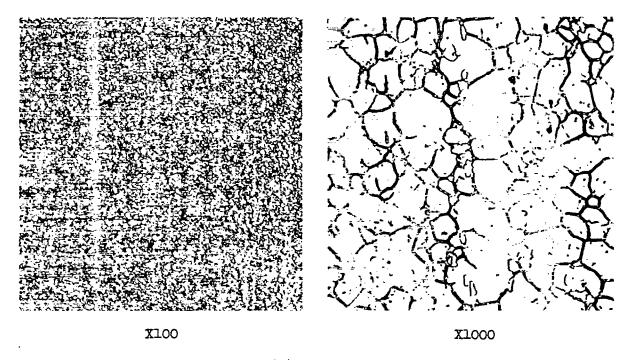


(b) Ruptured in 499 hours under 22,000 psi at 1350° F.

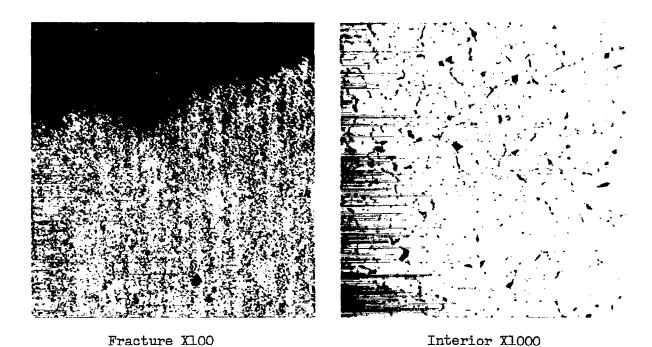
Figure 22.- Initial microstructure and microstructure after rupture testing of as-rolled and hot-cold-worked low-carbon N-155 alloy bar stock from heat 30276. Electrolytically etched in 10 percent chromic acid. (Treatment: As-rolled; 15-percent hot-cold-work at 1200° F.)

· ·

14



(a) Initial.



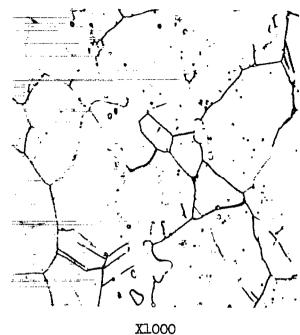
(b) Ruptured in 639 hours under 20,000 psi at 1350° F.

Figure 23.- Initial microstructure and microstructure after rupture testing for as-rolled and aged low-carbon N-155 alloy bar stock from heat 30276. Electrolytically etched in 10 percent chromic acid. (Treatment: As-rolled; 1350° F, 24 hr.)

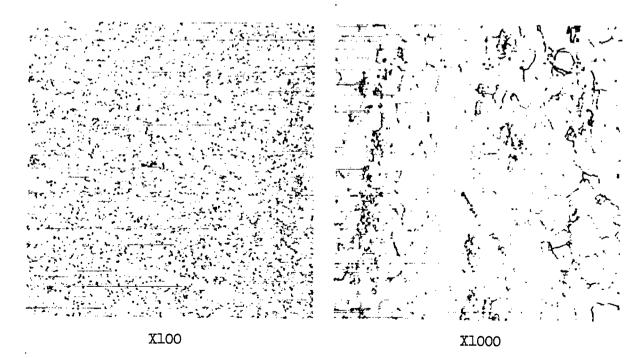
	•
	•
	•
	•

5

Xloo



(a) Heat Al726-initial.

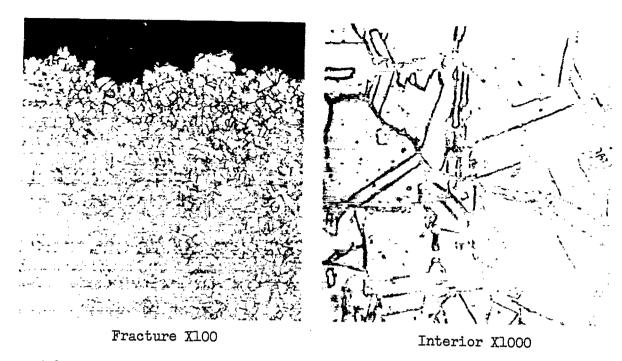


(b) Heat 30276-initial.

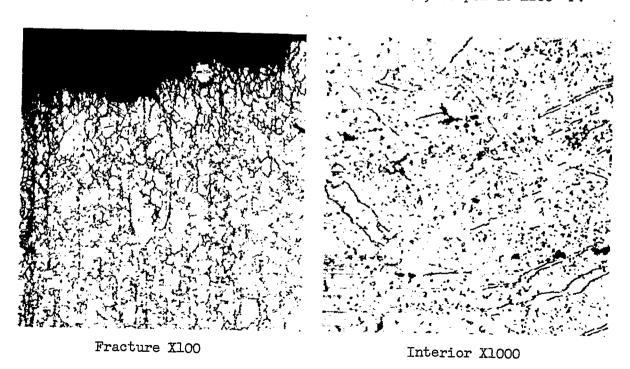
NACA __

Figure 24.- Initial microstructures and microstructures after rupture testing of solution-treated and hot-cold-worked low-carbon N-155 alloy bar stock from heats A1726 and 30276. Electrolytically etched in 10 percent chromic acid. (Treatment: 2050° F, 2 hr, water-quenched; 15-percent hot-cold-work at 1200° F.)

				:	
	•				
					-
					•



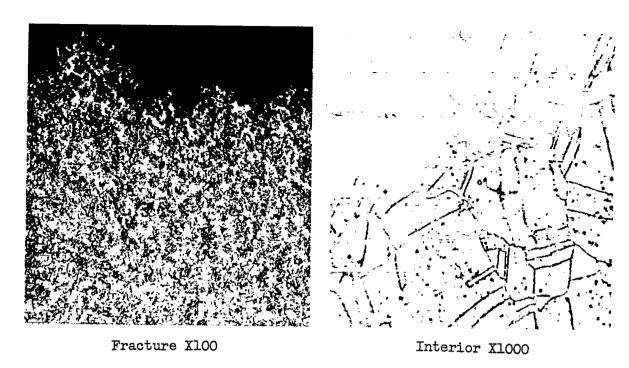
(c) Heat Al726 - ruptured in 942 hours under 48,000 psi at 12000 F.



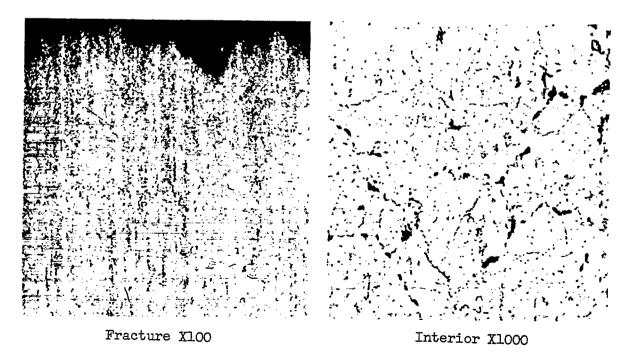
(d) Heat 30276 - ruptured in 1556 hours under 52,000 psi at 1200° F.

Figure 24.- Continued.

,			
			•
			4
		·	
			•



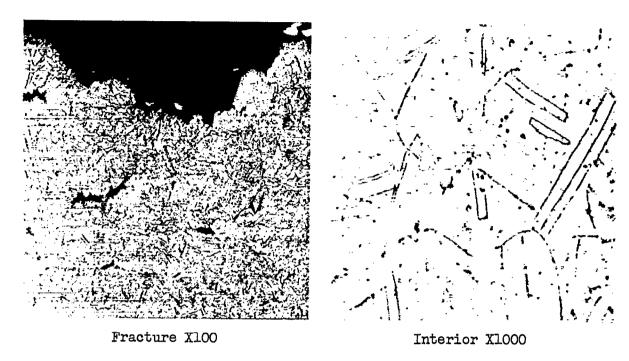
(e) Heat Al726 - ruptured in 56 μ hours under 33,000 psi at 13500 F.



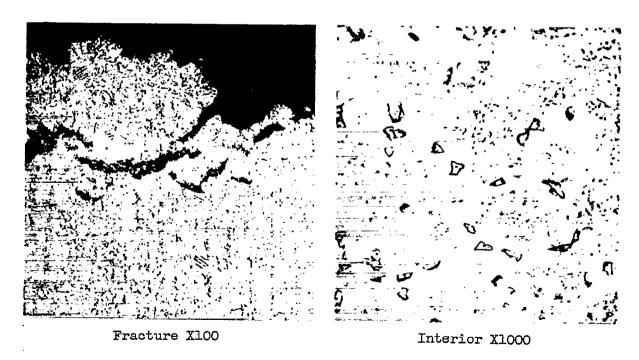
(f) Heat 30276 - ruptured in 593 hours under 30,000 psi at 1350° F.

Figure 24.- Continued.

		•
		•
•		
		•
		•
	,	
		•
		•

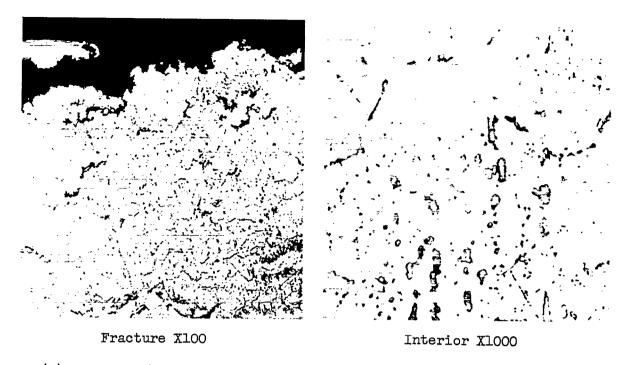


(g) Heat Al726 - ruptured in 739 hours under 18,000 psi at 1500° F.

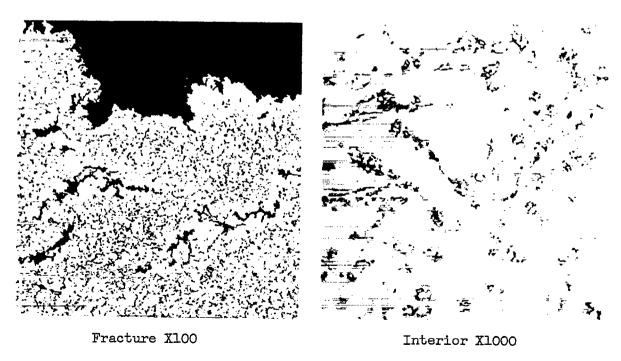


(h) Heat 30276 - ruptured in 421 hours under 13,000 psi at 1500° F. Figure 24.- Continued.

			•
			•
			•
			•
			,
			4
·			·

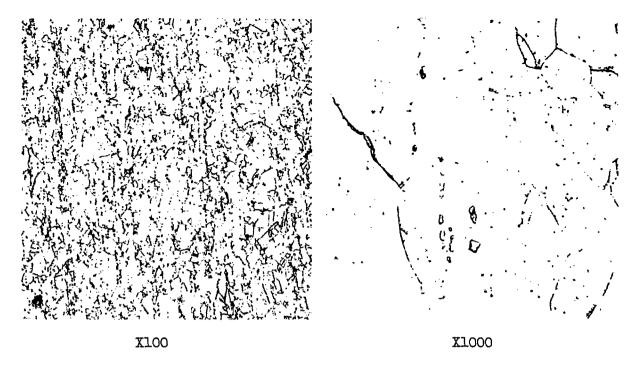


(i) Heat Al726 - ruptured in 676 hours under 7000 psi at 1650° F.

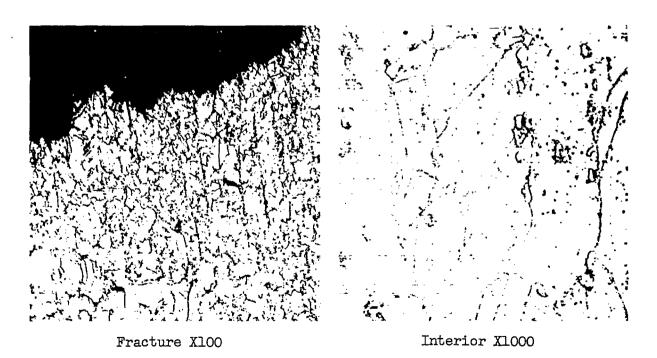


(j) Heat Al726 - ruptured in 1004 hours under 1850 psi at 1800° F.
Figure 24.- Concluded.

•			
			•
			~
•			•
			•
			~
			-
			F
			•



(a) Initial.

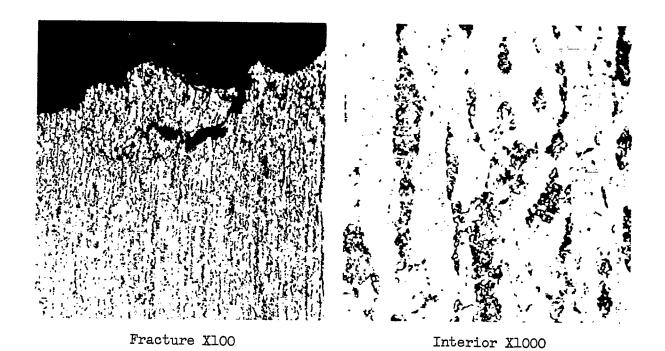


(b) Ruptured in 1142 hours under 55,000 psi at 1200° F.

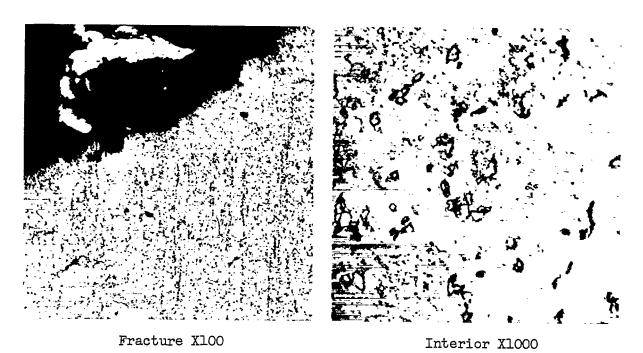
Figure 25.- Initial microstructure and microstructures after rupture testing of low-carbon N-155 alloy bar stock from heat 30276 hot-cold-worked 25 percent. Electrolytically etched in 10 percent chromic acid. (Treatment: 2050° F, 2 hr, water-quenched; 25-percent hot-cold-work at 1200° F.)

		•
		-
		•
		-

21

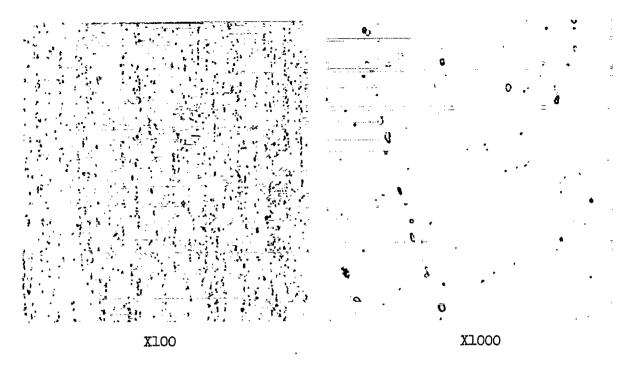


(c) Ruptured in 446 hours under 30,000 psi at 1350° F.

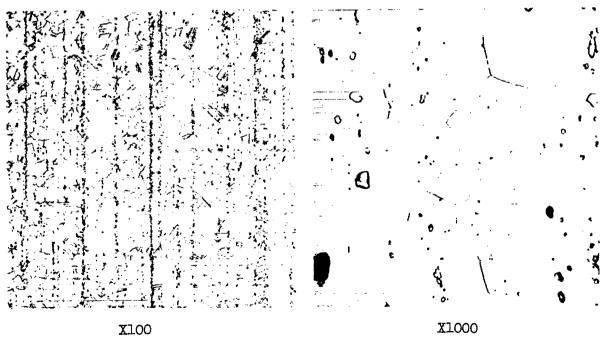


(d) Ruptured in 340 hours under 13,000 psi at 1500° F.
Figure 25.- Concluded.

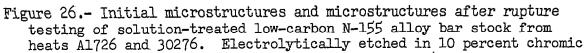
* *	4			
				~
				-
				•
				•
			i	
				•
				•
•				-
,				-
,				-
,				
,				-



(a) Heat Al726 - initial.



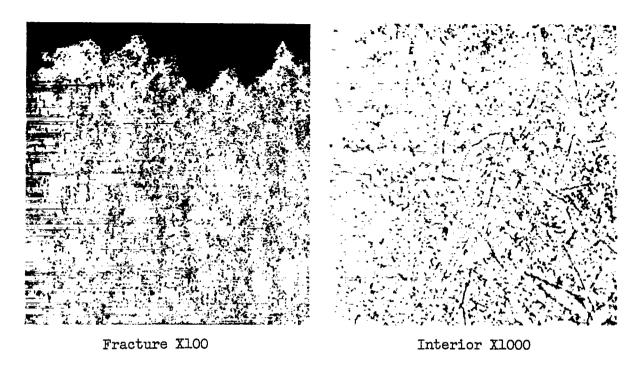
(b) Heat 30276 - initial.



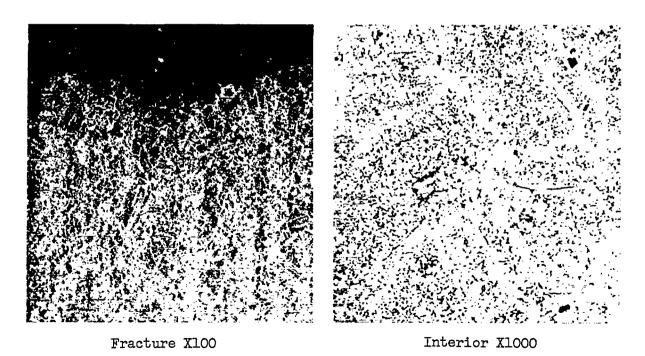
acid. (Treatment: 2100° F, 1 hr, water-quenched.)

		,	
		,	•
	·		
			•
			•
*			·

.3



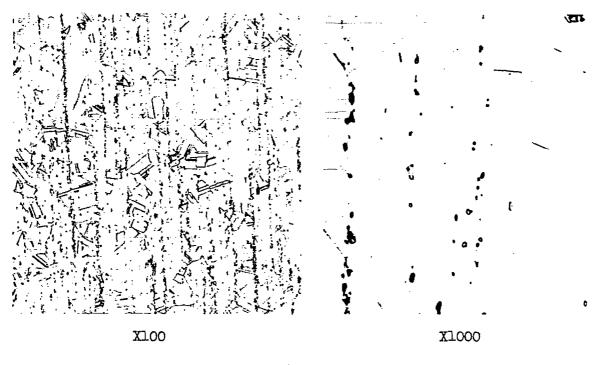
(c) Heat A1726 - ruptured in 696 hours under 25,000 psi at 1350° F.



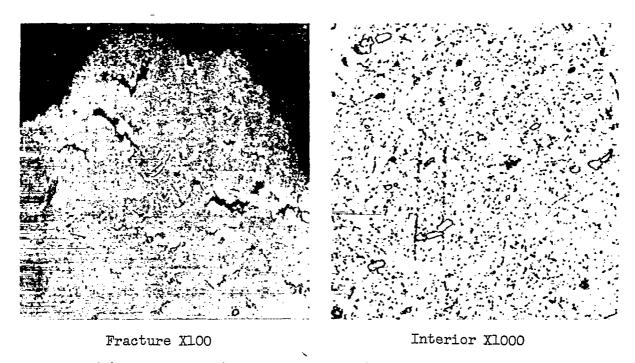
(d) Heat 30276 - ruptured in 997 hours under 22,000 psi at 1350° F.

Figure 26.- Concluded.

				•
		·		
				~
				-
			·	



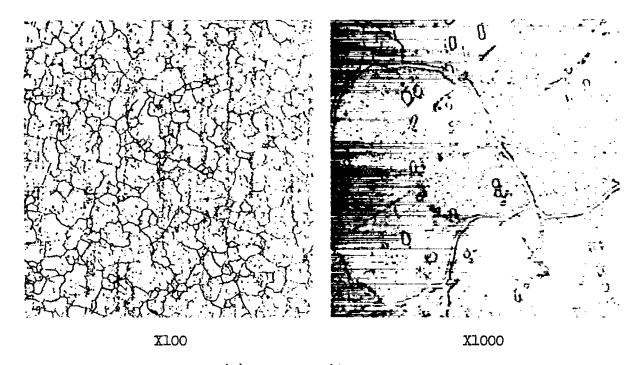
(a) Initial.



(b) Ruptured in 439 hours under 25,000 psi at 1350° F.

Figure 27.- Initial microstructure and microstructure after rupture testing of low-carbon N-155 alloy bar stock from heat 30276 solution-treated at 2200° F. Electrolytically etched in 10 percent chromic acid. (Treatment: 2200° F, 1 hr, water-quenched.)

			•
			•
			•
			•



(a) Heat Al726 - initial.

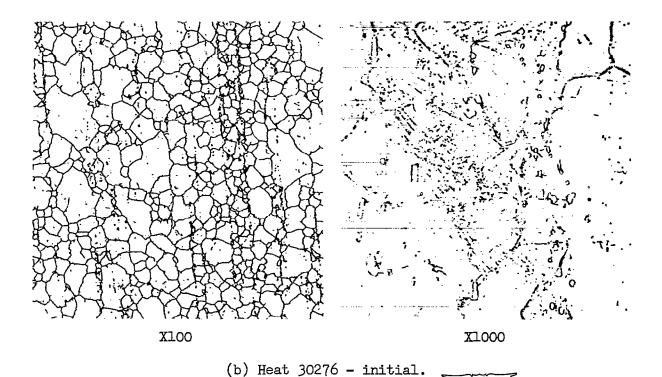
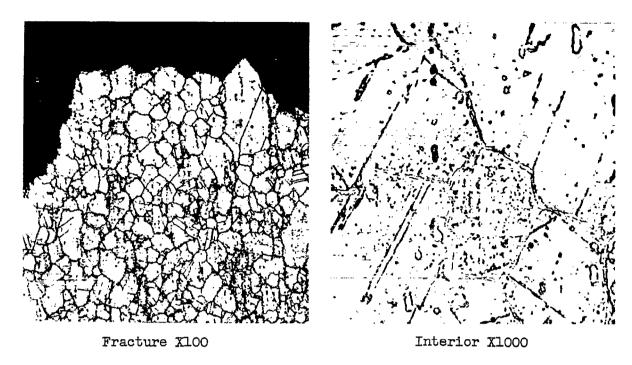
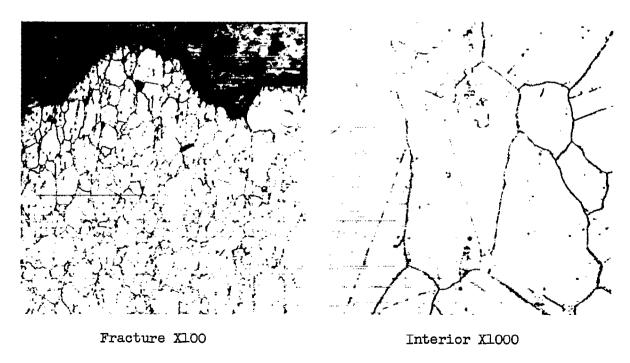


Figure 28.- Initial microstructures and microstructures after rupture testing of solution-treated and aged low-carbon N-155 alloy bar stock from heats A1726 and 30276. Electrolytically etched in 10 percent chromic acid. (Treatment: 2200° F, 1 hr, water-quenched; 1400° F, 24 hr.)

				•
				•
				•
				•
	•		·	



(c) Heat Al726 - ruptured in 446 hours under 44,000 psi at 1200° F.

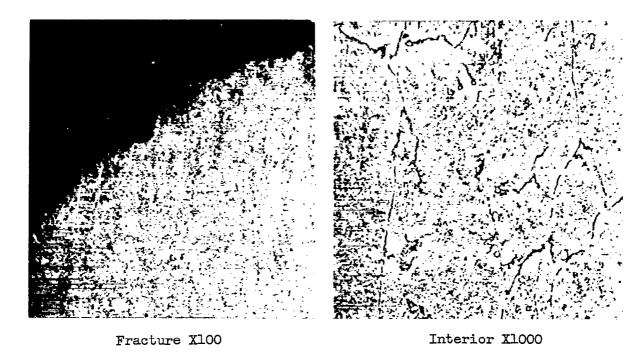


(d) Heat 30276 - ruptured in 133 hours under 47,000 psi at 1200° F.

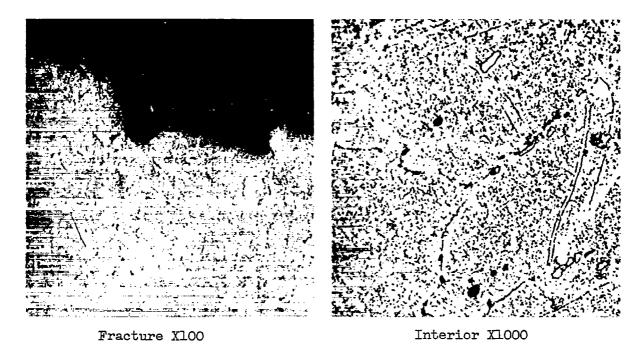
Figure 28.- Continued.

		E. S. C	` > *
			•
,			
			*

99



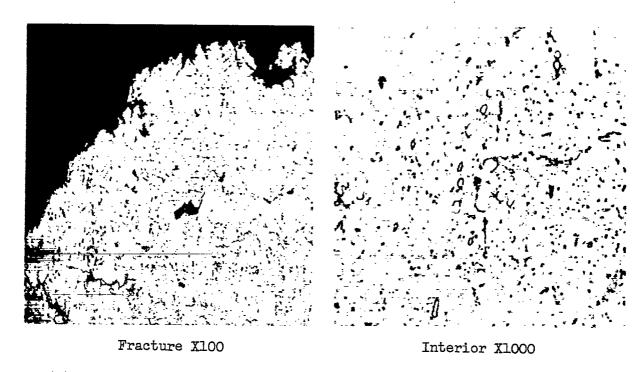
(e) Heat Al726 - ruptured in 441 hours under 26,000 psi at 1350° F.



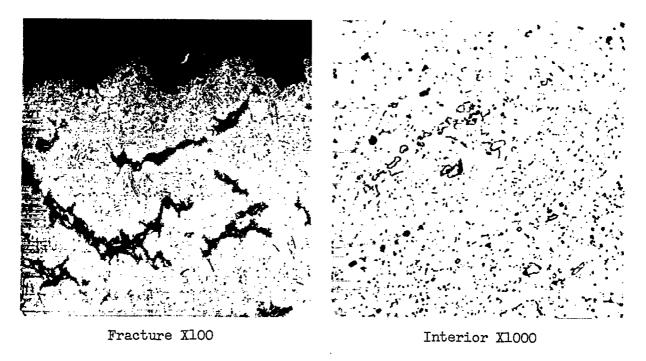
(f) Heat 30276 - ruptured in 726 hours under 25,000 psi at 1350° F. Figure 28.- Continued.

			•
			*
			•
			•
			·
			•

8

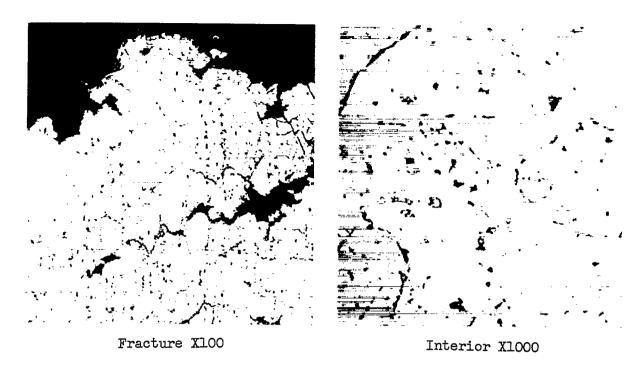


(g) Heat Al726 - ruptured in 1033 hours under 14,600 psi at 1500° F.

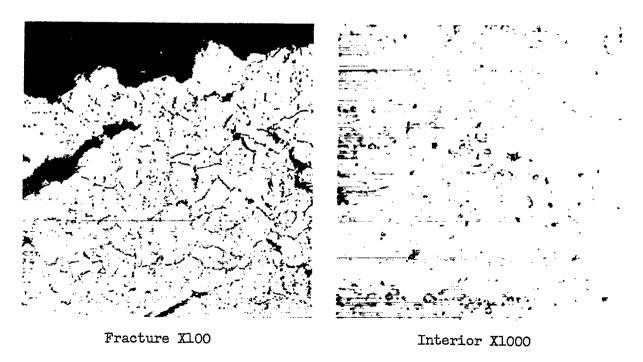


(h) Heat 30276 - ruptured in 1131 hours under 14,000 psi at 1500° F. Figure 28.- Continued.

* • en la maria de la companya de la co



(i) Heat Al726 - ruptured in 1384 hours under 7400 psi at 1650° ${\rm F.}$



(j) Heat A1726 - ruptured in 959 hours under 3300 psi at 18000 F.

Figure 28.- Concluded.

